

GenCore version 5.1.8
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protein - nucleic search, using frame_plus_p2n model

n on: May 31, 2006, 22:52:13 ; Search time 9316.63 Seconds

(without alignments)
5096.366 Million cell updates/sec

tle: US-10-048-116B-2

rfect score: 2660
quence: 1 MPCSRLILGLVALNTLSL.....VHEGLNHHHTKPSRTTCK 495

oring table:

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Ygapop 10.0 , Ygapext 0.5
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arched: 6366136 seqs, 31973710525 residues

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st-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	1924.5	72.3	1446	2	BD137962 Monovalen
3	1332	50.1	7528	2	AX080953 Sequence

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5	1324.5	49.8	1581	2	A78881	A78881 Sequence 1
6	1324.5	49.8	1581	6	MI7GHC2AA	X70423 M.musculus
7	1320.5	49.6	1341	2	I07390	I07390 Sequence 4
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9	1320.5	49.6	1570	6	AB097847	AB097847 Mus muscu
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ALIGNMENTS

RESULT 1	AX081280	Sequence 1	from Patent WO0109194.	1484 bp	DNA	linear	PAT 27-FEB-2001
LOCUS	AX081280	Sequence 1	from Patent WO0109194.	1484 bp	DNA	linear	PAT 27-FEB-2001
DEFINITION	AX081280	Sequence 1	from Patent WO0109194.	1484 bp	DNA	linear	PAT 27-FEB-2001
ACCESSION	AX081280	Sequence 1	from Patent WO0109194.	1484 bp	DNA	linear	PAT 27-FEB-2001
VERSION	AX081280.1	GI:13170129					
KEYWORDS		synthetic construct					
SOURCE		other sequences; artificial sequences.					
ORGANISM		1					
REFERENCE		Glaichenhaus, N. and Malherbe, L.					
AUTHORS		Recombinant proteins and molecular complexes derived therefrom,					
TITLE		analogous to molecules involved in immune responses					
JOURNAL		Patent: WO 0109194-A 1 08-FEB-2001;					
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10-048-116B-2 (1-495) x AX081280 (1-1484)

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RESULT 2

BD137962

LOCUS

DEFINITION

1446 bp DNA linear PAT 18-SEP-2002
 Monovalent MHC-binding domain fused proteins and conjugates,
 polyvalent MHC-binding domain fused proteins and conjugates,
 polymer MHC-binding domain fused proteins and conjugates, and
 utilization thereof.

ACCESSION

BD137962

VERSION

BD137962.1 GI:23232907

KEYWORDS

JP 2002504342-A/7.

SOURCE

synthetic construct

ORGANISM

other sequences; artificial sequences.

REFERENCE

1 (bases 1 to 1446)

Wucherpfennig, K.W. and Strominger, J.L.

Monovalent MHC-binding domain fused proteins and conjugates,

polyvalent MHC-binding domain fused proteins and conjugates,

polymer MHC-binding domain fused proteins and conjugates, and

utilization

Patent: JP 2002504342-A 7 12-FEB-2002;

PRESIDENT AND FELLOWS OF HARVARD COLLEGE

OS Artificial Sequence

PN JP 2002504342-A/7

PD 12-FEB-2002

PF 19-FEB-1999 JP 2000532537
 PR 19-FEB-1998 US 60/075351
 PI KAI W WUCHERPFENNIG, JACK L STROMINGER
 PC C12N15/09, A61K35/14, A61K47/48, C07K14/705, C07K16/00, C07K19/00,
 PC C12Q1/02;
 PC G01N33/53, C12N15/00
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 DEFINITION Sequence 3 from Patent WO0109303.
 ACCESSION AX080953
 VERSION AX080953.1 GI:13169890
 KEYWORDS
 SOURCE
 ORGANISM
 synthetic construct
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 other sequences; artificial sequences.
 REFERENCE
 AUTHORS
 TITLE
 Pit-3 ligand-encoding polynucleotide as a polynucleotide-based
 vaccine enhancer

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JOURNAL Patent: WO 0109303-A 3 08-FEB-2001;
VICAL INCORPORATED (US)
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LOCUS
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ACCESSION CO897414
VERSION CO897414.1 GI:55582199
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE
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AUTHORS other sequences; artificial sequences.
TITLE Immune response; antigen presentation; T cell activation; T cell
JOURNAL Patent: WO 2004091655-A 1 28-OCT-2004;
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VERSION	A78881.1	GI:6090456	
KEYWORDS	unidentified		
SOURCE	unclassified sequences.		
ORGANISM	1 (bases 1 to 1581)		
REFERENCE	Boulain,J. and Ducancel,P.		
AUTHORS	HYBRID PROTEIN COMPLEXES, THEIR PROCESS OF PREPARATION, AND THEIR		
TITLE	APPLICATIONS AS AN AGENT IN DIAGNOSTIC AND IN THE THERAPEUTIC FIELD		
JOURNAL	OR AS A REAGENT RELEVANT IN MEDICAL APPROACHES		
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>  Okawa, H., Nakata, M. and Yuasa, Y.
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>  KANKYO MENEKI GIJUTSU KENKUSYO KK
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 ACCESSION AB097847
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 REFERENCE
 AUTHORS Nishi K., Imajuku Y., Nakata M., Ohde K., Miyake S., Morimune K., Kawata M. and Ohkawa H.
 TITLE Molecular characteristics of the monoclonal and recombinant antibodies specific to the insecticide malathion
 JOURNAL Unpublished
 REFERENCE
 AUTHORS Nishi K., Imajuku Y., Nakata M., Ohde K., Miyake S., Morimune K., Kawata M. and Ohkawa H.
 TITLE Direct Submission
 JOURNAL Submitted (11-DEC-2002) Kosuke Nishi, Kobe University, Research Center for Environmental Genomics; 1-1 Rokkodai-cho, Nada-ku, Kobe, Hyogo 657-8501, Japan (E-mail: nishikosuke@yahoo.co.jp, Tel:81-78-803-5863, Fax:81-78-871-3617)
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693  -TGCAAATGTGCCCCCCCAGCAGACCAAGGTGGACAGAAGAAAAT----- 740
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 Gaps: 13

-10-048-116B-2 (1-495) x CS125905 (1-1407)

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238 GGTACTAATCAATGAGAAGTTCAAGAGCAAGGCCACACTGACTGTAGACAAATCCTCC 297
66 LysLysThrValTyrArgLeuPro----- 73
298 AGCAGAGCTACATGAGCTCAGAGCCTGACATCTGAGGACTCTGCGGTCTATTATTGT 357
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358 GBACTGGGACAG-----GGCTACTGGGGCCAGGCACACTAGTCACCGCTCTCTCAGCC 411
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412 AAA-----ACAAACAGCC 423
114 ProGlnAlaThrValPheProLysSerProVal-----LeuLeuGlyGlnProAsn 130
424 CCA-----TCGGTCTATCCATCGCCCTGTGTGGAGATACAACTGGCTCTCGGTG 477
131 ThrLeuIleCysPheValAspAsnIlePheProProValIleAsnIleThrTrpLeuArg 150
478 ACTTAGGATGCTGGTCAAGGGTATTTCCTGAGCCAGTACCTTGACCTGG----- 531
151 AsnSerLysSerValThrAspGlyValTyrGluThrSerPheLeuValAsnArgAspHis 170
532 AACTCTGGATCCCTGTCAGTGGTGTGCACACCTTCCAGAGTCTCTGAGCTGAC--- 588
171 SerPheHisLysLeuSerTyrLeuThrPheIleProSerAspAspAspIleTyrAspCys 190
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618 AACCTGAGCAGCTGG----- 633
210 oAlaProMetSerGluLeuThrGluThrGlyGlyGlySerThrThrAlaProSerAl 230
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700 -----GAGCCAGAGGGCCCAATCA 722
270 eProCysProProCysLysCysProAlaProAsnLeuLeuGlyGlyProSerValPheI 290
723 GCCCTGTCTCCATGCAATGCCAGCACCTAACTCTCTGGTGGCCCATCGCTTCTAT 782
290 ePheProProLysIleLysAspValLeuMetIleSerLeuSerProIleValThrCysVa 310
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310 lValValAspValSerGluAspAspProAspValGlnIleSerTrpPheValAsnAsnVa 330

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903 GGAAGTACACACAGCTCAGACACAAACCCATAGAGAGGATTACAACAGTACTCTCCGGGT 962
350 lValSerAlaLeuProIleGlnHisGlnAspTrpMetSerGlyLysGluPheLysCysLys 370
963 GGTGAGTGGCTCCCTCCCATCCAGCACAGGACTGGATGAGTGGCAGGAGTTCAATGCAA 1022
370 sValAsnAsnLysAspLeuProAlaProIleGluArgThrIleSerLysProLysGlySe 390
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390 sValArgAlaProGlnValTyrValLeuProProGluGluGluMetThrLysLysG 410
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490 rArgThrProGlyLys 495
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LOCUS             Sequence 49 from Patent WO2005061544.
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ACCESSION         CS126190.1  GI:71059163
VERSION           Mus musculus (house mouse)
KEYWORDS          Mus musculus
SOURCE            Mus musculus
ORGANISM          Mus musculus
REFERENCE         Ellis,J.H.
AUTHORS           Nogo-a neutralising immunoglobulins for treatment of neurological
TITLE             diseases
JOURNAL           Patent: WO 2005061544-A 49 07-JUL-2005;
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Pred. No.: 6,08e-123 Length: 1407
Score: 1320.00 Matches: 285
Percent Similarity: 64.0% Conservative: 26
Best Local Similarity: 58.6% Mismatches: 68
Query Match: 49.6% Indels: 108
DB: 2 Gaps: 13

US-10-048-116B-2 (1-495) x CS126190 (1-1407)

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66 LysLysThrValTyrArgLeuPro----- 73
298 AGCAGAGCTATGCACTGAGAGCTGAGAGCTCTGAGAGCTCTGCGGTCTATTATTGT 357
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478 ACTTAGATGCTGGTCAAGGGTTATTTCTCTGAGCCAGTGACCTTGACCTGG----- 531
151 AenSerLysSerValThrAaspGlyValTyrGluThrSerPheLeuValAenAaspHis 170
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171 SerPheHisLysLeuSerTyrLeuThrPheIleProSerAaspAaspIleTyrAaspCys 190
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191 LysValGluHis-TrpGlyLeuGluGluProValLeuLysHisTrpGluProGluIlePr 210
618 AACCTCGAGCACCTGG----- 633
210 oAlaProMetSerGluLeuThrGluThrGlyGlyGlySerThrThrAlaProSerAl 230
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691 CAAGGTGGACAGAGAAATTT----- 699
250 uGlnAlaLeuGluLysGluLeuAlaGlnAlaSerGluProArgGlyProThrIleLy 270
700 -----GAGCCCAAGAGGCCCAACATCAA 722
270 aProCysProProCysLysCysProAlaProAenLeuLeuGlyGlyProSerValPheIl 290
723 GCCTGTCTCTCATGCAATGCCAGCACCTTAACCTCTCTGGTGGCCCATCGTCTTCAT 782
290 ePheProProLysIleLysAaspValLeuMetIleSerLeuSerProIleValThrCysVa 310
783 CTTCCTCCAAAGATCAAGAGTGTACTCATGATCTCTCTGAGCCCAATGATCATGTGT 842
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963 GGTGAGTGCCTCTCCCATCCAGCACAGGACTGGATGATGGCAAGGAGTTCAATGCAA 1022
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Qy 410 nValThrLeuThrCysMetValThrAaspPheMetProGluAaspIleTyrValGluTrpTh 430
Db 1143 GGTCACTCTGACCTGTCATGTCACAGACTTCATGCTGAAGACATTTACGTGGAGTGGAC 1202
Qy 430 rAenAenGlyLysThrGluLeuAenTyrLysAenThrGluProValLeuAaspSerAaspG1 450
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Qy 490 rArgThrProGlyLys 495
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RESULT 12
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LOCUS Sequence 1 from Patent WO2005068503.
DEFINITION CS138860
ACCESSION CS138860.1 GI:73530223
VERSION CS138860.1
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muroidea; Muridae; Murinae; Mus.

REFERENCE
1 Liu, C.
AUTHORS M-csf-specific monoclonal antibody and uses thereof
TITLE Patent: WO 2005068503-A 1 28-JUL-2005;
JOURNAL CHIRON CORPORATION (US); Liu, Cheng (US)
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Location/Qualifiers
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 MMIGG6
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 linear
 ROD 17-NOV-2004
 Mouse mRNA for gamma-2a-immunoglobulin heavy-chain.
 'SSION
 V00798
 V00798.1
 GI:51835

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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 1095)
Skorav,J.L., Auffray,C. and Rougeon,F.
Structure of the constant and 3' untranslated regions of the murine
Balb/c gamma 2a heavy chain messenger RNA
Nucleic Acids Res. 8 (14), 3143-3155 (1980)
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Copyright (c) 1993 - 2006 Bioceleration Ltd.

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US-10-048-116B-2

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Maximum Match 100%

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3: Geneseqn2000s.*
4: Geneseqn2001as.*
5: Geneseqn2001bs.*
6: Geneseqn2002bs.*
7: Geneseqn2002bs.*
8: Geneseqn2003bs.*
9: Geneseqn2003bs.*
10: Geneseqn2003cs.*
11: Geneseqn2003ds.*
12: Geneseqn2004as.*
13: Geneseqn2004bs.*
14: Geneseqn2005s.*
15: Geneseqn2006s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

sult No.	Score	Query Match	Length	ID	Description
1	2655	99.8	1484	5 AAF55098	Aaf55098 DNA encod
2	2247	84.5	1676	4 ABI99041	Abi99041 Murine pc
3	1924.5	72.3	1446	2 AAT99707	Aat99707 DR2-IGG f

4	1924.5	72.3	1446	2	AAX87813	Aax87813 HLA-DR2 a
5	1924.5	72.3	1446	14	ADW44282	Adw44282 DR2-IGG f
6	1738	65.3	2346	4	ABI99027	Abi99027 IAS MBP 1
7	1737.5	65.3	2343	4	ABI99033	Abi99033 MBP 90-10
8	1521	57.2	2053	4	ABI99029	Abi99029 IAS MBP 9
9	1521	57.2	2059	4	ABI99032	Abi99032 MBP 1-14
10	1332	50.1	7528	4	AAF30316	Aaf30316 Biciatron
11	1328.5	49.9	3973	13	ADT77690	Adt77690 Monoclona
12	1324.5	49.8	1581	2	AAQ48037	Aaq48037 Monoclona
13	1321.5	49.7	1560	14	AED19725	Aed19725 Anti-PrP
14	1320	49.6	1407	14	AEB21727	Aeb21727 Anti-Nogo
15	1320	49.6	1407	14	AEB08761	Aeb08761 antibody
16	1319.5	49.6	1401	14	AEC20762	Aec20762 M-CSF spe
17	1318.5	49.6	990	12	ADL15694	Adl15694 Murine im
18	1318.5	49.6	1569	14	ADV26108	Adv26108 Mouse OKT
19	1318.5	49.6	1569	14	ADW71834	Adw71834 Murine OK
20	1318.5	49.6	1570	2	AAQ12637	Aaq12637 Monoclona
21	1318.5	49.6	1570	12	ADQ91058	Adq91058 Murine OK
22	1317.5	49.5	6729	4	AAF30341	Aaf30341 Biciatron
23	1315.5	49.5	1645	2	AAQ54652	Aaq54652 T84.12 He
24	1312.5	49.3	1341	1	AAQ91859	Aaq91859 Chimeric
25	1309.5	49.2	1356	12	ADN97544	Adn97544 Artificia
26	1300.5	48.9	1389	14	AEA27480	Aea27480 Monoclona
27	1285.5	48.3	1140	10	ADE85817	Ade85817 Murine in
28	1284	48.3	1194	2	AAV55415	Aav55415 Chimeric
29	1281.5	48.2	1524	14	AEE21946	Aee21946 Single ch
30	1280.5	48.1	1473	13	ADS31748	Ads31748 DNA encod
31	1280.5	48.1	1473	13	ADS92750	Ads92750 DNA encod
32	1280	48.1	1045	12	ADO07566	Ado07566 Fusion pr
33	1280	48.1	1509	14	AEE21942	Aee21942 Single ch
34	1279.5	48.1	1140	10	ADE85819	Ade85819 Murine mu
35	1278	48.0	1131	2	AAV55416	Aav55416 Chimeric
36	1275	47.9	1108	12	ADO07578	Ado07578 Fusion pr
37	1275	47.9	1108	12	ADO07577	Ado07577 Fusion pr
38	1275	47.9	1461	6	AAD22972	Aad22972 Mouse Zal
39	1272	47.8	729	3	AAZ35704	Aaz35704 Human gly
40	1270	47.7	1707	3	AAZ35706	Aaz35706 Human gly
41	1269.5	47.7	1431	14	AEB12362	Aeb12362 Fusion pr
42	1269.5	47.7	1431	14	AED64242	Aed64242 hOGH-mFC-
43	1269	47.7	1188	2	AAT59349	Aat59349 1-153 c-m
44	1269	47.7	1275	2	AAT62850	Aat62850 Mouse sol
45	1268	47.7	699	3	AAZ51300	Aaz51300 Murine im

ALIGNMENTS

RESULT 1

AAF55098

ID AAF55098 standard; DNA; 1484 BP.

AC AAF55098;

DT 15-MAY-2001 (first entry)

DE DNA encoding a fusion protein comprising an alpha chain of MHC.

KW Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
major histocompatibility complex; Fc region; antigen; T lymphocyte;
immunostimulant; vaccine; infection; tumour; ss.

OS Synthetic.

XX Key Location/Qualifiers

FT CDS 1..1482

FT /*tag= a

XX WO200109194-A1.

XX 08-FEB-2001.

XX 28-JUL-2000; 2000WO-FR002193.

XX 29-JUL-1999; 99FR-00009862.

(CNRS) CNRS CENT NAT RECH SCI.

Glaichenhaus N, Malherbe L;

WPI; 2001-182944/18.

P-PSDB; AAB67480.

New soluble recombinant protein, useful e.g. as immunostimulant, comprises dimeric major histocompatibility complex molecule fused to immunoglobulin Fc region.

Example 1; Page 31-33; 43pp; French.

The specification describes soluble recombinant proteins that comprise at least a dimer formed from the alpha and beta-chains of MHC (major histocompatibility complex) Class I and II molecules in which at least one chain has, attached to its C-terminus, at least part of the Fc region of an immunoglobulin. The recombinant proteins, when linked to an antigenic peptide, are used to count and/or purify antigen-reactive T lymphocytes and to characterize their phenotype, e.g. in preclinical evaluation of vaccines. They are also used as immunostimulants, particularly for vaccine development (against infections and tumours), to count and determine phenotype of autoreactive T cells in subjects with, or at risk of developing, autoimmune diseases, e.g. for staging or evaluating treatments, and to purify and/or enrich Ag-reactive T cells from cell cultures or patient samples, for use in subsequent curative or preventative cellular therapy. The present sequence encodes a recombinant protein of the invention, comprising an alpha chain of MHC molecules

Sequence 1484 BP; 414 A; 394 C; 362 G; 314 T; 0 U; 0 Other;

gment Scores:

cl. No.:	1.06e-218	Length:	1484
re:	2655.00	Matches:	494
cent Similarity:	100.0%	Conservative:	0
t Local Similarity:	100.0%	Mismatches:	0
ry Match:	99.8%	Indels:	0
	5	Gaps:	0

10-048-116B-2 (1-495) x AAF55098 (1-1484)

```

1 MetProCysSerArgAlaLeuLeuGlyValLeuAlaLeuAenThrMetLeuSerLeu 20
1 ATCCGTCGACGAGAGCTGATTCGGGGTCTCGCCCTCGAACCACCATGTCAGCCTC 60
21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrValTyr 40
61 TCCGAGGTGAAGACGACATGAGGCCGACCACGTCAGGCTTCTATGGTACAACTGTTTAT 120
41 GlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
121 CAGTCTCTCGAGACATTTGGCCAGTACACACATGATTTGATGGTATGATGTTCTAT 180
61 ValAspLeuAspLysGlyLysThrValTrpArgLeuProGluPheGlyGlnLeuLeu 80
181 GTGGACTTGGATAAGAGAAAACCTGCTGGAGGCTTCTGAGTTGGCCAATTGATATC 240
81 PheGluProGlnGlyGlyGlnAnIleAlaGluLysHisAsnLeuGlyIleLeu 100
241 TTTGAGCCCCAAGGTGGATCGCAAAACATAGCTGCAGAAAACACAACTTGGGAATCTTG 300
101 ThrLysArgSerAspPheThrProAlaThrAenGluAlaProGlnAlaThrValPhePro 120
301 ACTAAGAGTCAAAATTCACCCAGCTACCAATGAGGCTCTCAAGGACATGTGTCCCC 360
121 LysSerProValLeuLeuGlyGlnProAenThrLeuIleCysPheValAspAsnIlePhe 140
361 AAGTCCCTGTGCTGCTGGGTGAGGCCAACACCCCTTATCTGCTTTGTGGACAAATCTTC 420
141 ProProValIleAsnIleThrTrpLeuArgAenSerLysSerValThrAspGlyValTyr 160
421 CCACCTGTGATCAACATCACATGGCTCGAAATAGCAAGTCAGTCAGCAGCGGGTTTAT 480

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RESULT 2

ABI99041

ID ABI99041 standard; cDNA; 1676 BP.

XX

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Qy 161 GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe 180
Db 481 GAGACCAGAGCTTCCTGGTCAACCGTGACCATTCCTTCCACAAGCTGTCTTATCTCACCTTC 540
Qy 181 IleProSerAspAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGluGluPro 200
Db 541 ATCCCTTCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 600
Qy 201 ValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGluThrGly 220
Db 601 GTTCTGAACACTGGGAACCTGAGATTCAGACCCCATGTGTCAGAGCTGACAGAACTGGA 660
Qy 221 GlyGlyGlySerThrThrAlaProSerAlaGlnLeuGlyGluValGluGlnAlaLeuGlu 240
Db 661 GETGGAGATCCACTACAGCTCCATCAGCTCAGCTCGAAAAAGAGCTCCAGGCCCTGGAG 720
Qy 241 LysGluAsnAlaGlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeuAlaGlnAla 260
Db 721 AAGGAAAATGCACAGCTGGAAATGGGAGTTGCAAGCACTGGAAAAAGGAACCTGGCTCAGGCA 780
Qy 261 AlaSerGluProArgGlyProThrIleLysProCysProProCysLysCysProAlaPro 280
Db 781 GCATCTGAGCCGACAGAGGCCCAATCAAGCCCTGTCTCTCATGCAATATGCCACGACCT 840
Qy 281 AsnLeuLeuGlyGlyProSerValPheIlePheProProLysIleLysAspValLeuMet 300
Db 841 AACCTCTGGGTGGACCATCGTCTTCTATCTTCCCTCCAAAGATCAAGGATGTACTCATG 900
Qy 301 IleSerLeuSerProIleValThrCysValValValAspValSerGluAspAspProAsp 320
Db 901 ATCTCCTGAGCCCATAGTCACATGTGTGGTGGTGTGAGCGAGGATGAGCCCAAGAT 960
Qy 321 ValGlnIleSerTrpPheValAsnAsnValGluValHisThrAlaGlnThrGlnThrHis 340
Db 961 GTCCAGATCAGCTGGTTGTGAAACAGCTGGAAGTACACACAGCTCAGACACAACCCAT 1020
Qy 341 ArgGluAspTyrAenSerThrLeuArgValValSerAlaLeuProIleGlnHisGlnAsp 360
Db 1021 AGAGAGGATTACAAACAGTACTCTCCGGGTGGTGTGAGTGCCTCCCTCCATCCAGCAGGAC 1080
Qy 361 TrpMetSerGlyLysGluPheLysCysLysValAsnAsnLysAspLeuProAlaProIle 380
Db 1081 TGGATGATGGCAGGAGGTTCNAATGCAAGGTCAACAAACAAAGACCTCCAGCGCCCATC 1140
Qy 381 GluArgThrIleSerLysProLysGlySerValArgAlaProGlnValTyrValLeuPro 400
Db 1141 GAGAGAACCATCTCANAAACCCAAAGGTCAGTAAGAGCTCCACAGGTATATGTCTTGCT 1200
Qy 401 ProProGluGluMetThrLysLysGlnValThrLeuThrCysMetValThrAspPhe 420
Db 1201 CCACCAAGAAGAGATGAGTCAAGAAAACAGGTCACTCTGACCTGCATGGTGCACAGACTTC 1260
Qy 421 MetProGluAspIleTyrValGluTrpThrAsnAsnGlyLysThrGluLeuAsnTyrLys 440
Db 1261 ATGCCTCAAGACATATTCAGTGGAGTGGACCAACCAACGGGAAAACAGAGCTAAACTACAG 1320
Qy 441 AsnThrGluProValLeuAspSerAspGlySerTyrPheMetTyrSerLysLeuArgVal 460
Db 1321 AACACTGACAGCTCTGGACTCTGATGGTTCTTACTTTCATGTACAGCAAGCTGAGAGTG 1380
Qy 461 GluLysLysAsnTrpValGluArgAsnSerTyrSerCysSerValValHisGluGlyLeu 480
Db 1381 GAAAGAAGAACTGGGTGGAAAAGAAATAGCTACTCTCTGTTTCAGTGGTGCACAGAGGCTG 1440
Qy 481 HisAsnHisThrThrLysSerPheSerArgThrProGly 494
Db 1441 CACAATCACCACACGACTAAGAGCTTCTCCGGGACTCCGGGT 1482

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1222 CTCGGGTGGTCAAGTCCCTCCCATCCAGCAGCAGTGGATGATGAGTGGCAAGGAGTTC 1281
368 LysCysValAsnAspLeuProAlaProIleGluArgThrIleSerIysPro 387
1282 AAATGCAAGGTCAACAACAAGACCTCCAGCGGCCCATCGAGAGAACCATCTCAAAACCC 1341
388 LysGlySerValArgAlaProGlnValTyrValLeuProProGluGluGluMetThr 407
1342 AAAGGTCAGTAAGAGCTCCACAGGTATATGTTGCTCCACCAAGAGAGAGATGACT 1401
408 LysLysGlnValThrLeuThrCysMetValThrAspPheMetProGluAspIleTyrVal 427
1402 AAGAAACAGGTCACTCTGACCTGCATGCTGCATGCTGCATGCTGCATGCTGCATGCTG 1461
428 GluTyrThrAsnAsnGlyValThrGluLeuAsnTyrLysAsnThrGluProValLeuAsp 447
1462 GAGTGGACCAACAACGGGAACAGAGCTAACTACAAGAACTGAACACCTCTGGAC 1521
448 SerAspGlySerTyrPheMetTyrSerLysLeuArgValGluLysAsnTyrValGlu 467
1522 TCTGATGTTCTTACTTCTATGATGACCAAGCTGAGAGTGGAAAGAGAACTGGGTGAA 1581
468 ArgAsnSerTyrSerCysSerValValHisGluGlyLeuHisAsnHisThrThrLys 487
1582 AGAAATAGCTACTCTCTGTTCAAGTGTCCACAGAGGTCTTGCAATCAACACAGCTAAG 1641
488 SerPheSerArgThrProGlyLys 495
1642 AGCTTCTCCCGACTCCGGGTAAA 1665

UTL 3

99707

AAT99707 standard; cDNA; 1446 BP.

AAT99707;

17-OCT-2003 (revised)

17-AUG-1998 (first entry)

DR2-IgG fusion construct.

Major histocompatibility complex class II; MHC class II; human; mouse;
fusion protein; HLA-DR2; DR2*0101; binding domain; Fos;
dimerization domain; IgG; allergy; autoimmune disease; vaccine;
multiple sclerosis; therapy; ss.

Homo sapiens.

Mus musculus.

Chimeric.

W09806749-A2.

19-FEB-1998.

15-AUG-1997; 97WO-US014503.

16-AUG-1996; 96US-0024077P.

(HARD) HARVARD COLLEGE.

Wucherpennig KW, Strominger JL;

WPI; 1998-159459/14.

New Class II MHC fusion proteins - comprising a MHC Class II binding
domain and a dimerization domain or an immunoglobulin region used for
modulating immune responses.

Example; Page 49; 76pp; English.

This nucleotide sequences codes for a bivalent DR2 fusion protein
obtained by fusion of the Fc portion of IgG2a to the 3' end of a DR-alpha
-Fos cDNA construct (see AAV16866). The Fc portion was amplified by RT-

CC PCR from mouse hybridoma L243. The PCR product was then fused in frame
CC with the DR-alpha-Fos construct by overlapping PCR. The DR2-IgG fusion
CC was expressed in the Drosophila Schneider cell system. The invention
CC relates to new soluble monovalent and multivalent Class II MHC fusion
CC proteins comprising a MHC Class II binding domain and a dimerization
CC domain or an immunoglobulin region that can be used for the treatment
CC of allergic and autoimmune diseases (e.g. multiple sclerosis), for
CC transplanting a subject to foreign tissue before or after organ or tissue
CC transplantation, or for vaccination against pathogens. (Updated on 17-OCT
CC -2003 to standardise OS field)

XX Sequence 1446 BP; 414 A; 375 C; 356 G; 301 T; 0 U; 0 Other;
SQ

Alignment Scores:

Pred. No.:	8.16e-156	Length:	1446
Score:	1924.50	Matches:	364
Percent Similarity:	83.2%	Conservative:	32
Best Local Similarity:	76.5%	Mismatches:	73
Query Match:	72.3%	Indels:	7
DB:	2	Gaps:	4

US-10-048-116B-2 (1-495) x AAT99707 (1-1446)

Qy 26 AspleGluAlaAspHisValGlyPheTyrGlyThrThrValTyrGlnSerProGlyAsp 45

Db 13 GAGATCAAGAAGAACATGTG---ATCATCCAGCGCGAGTTCTATCTGAAATCCCTGCCAA 69

Qy 46 IleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyrValAspLeuAspLys 65

Db 70 TCAGGCGAGTTTATGTTTGACTTTGATGGTGTGATGAGATTTTCCATGTGGATATGGCAAG 129

Qy 66 LysLysThrValTyrArgLeuProGluPheGlyGlnLeuLeuPheGluProGlnGly 85

Db 130 AAGGAGACGCTCTGGCGCTTGAAGAAATTTGGACGATTTCAGCTTTGAGGCTCAAGGT 189

Qy 86 GlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGlyIleLeuThrLysArgSerAsn 105

Db 190 GCATTGGCCAAACATAGCTGTGGCAAAAGCCCAACTTGGAAATCATGACAAAGCGCTCCAAC 249

Qy 106 PheThrProAlaThrAsnGluAlaProGlnAlaThrValPheProLysSerProValLeu 125

Db 250 TATACTCCGATCCCAATGTACCTCCAGAGTAACTGTGCTCAGAACAGCCCTGTGGAA 309

Qy 126 LeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePheProProValIleAsn 145

Db 310 CTGAGAGAGCCCAACGCTCTCATCTGTTTCATAGACAAAGTTCAACCCACAGTGGTCAAT 369

Qy 146 IleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyrGluThrSerPheLeu 165

Db 370 GTCACGTGGCTTCGAAATGGAAACCTGTCAACACAGAGGTGTCAGACAGTCTTCTCTG 429

Qy 166 ValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPheIleProSerAspAsp 185

Db 430 CCAGGGAAGACCACTTTTCGCAAGTTTCACTATCTCCCTTCCTGCCCTCACTGAG 489

Qy 186 AspleTyrAspCysLysValGluHisTrpGlyLeuGluProValLeuLysHisTrp 205

Db 490 GACGTTTACGACTGACGGGTGGAGCACTGGGGCTTGGATGAGCTCTTCTCAAGCACTGG 549

Qy 206 GluProGluLeuProAlaProMetSerGluLeuThrGluThr---GlyGlyGlySer 224

Db 550 GAGTTTGTGCTCCAGCCCTCTCCAGAGACTACAGAGGTGCGAGGTGGCGGGCGGT 609

Qy 225 ThrThr-----AlaProSerAlaGlnLeuGluLysGluLeuGlnAlaLeuGlu 240

Db 610 TTAACGTATACATCCAGCGGAGACAGATCACTTGAACAGAGAGTCTGCGTTCGAG 669

Qy 241 LysGluAsnAlaGlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeu---AlaGln 259

Db 670 ACCGAGATTGCCAATCTACTGAAAGAGAGAGAACTGGAGTTTCATCTGGCGGCCCAT 729

Qy 260 AlaAlaSerGluProArgGlyProThrIleLysProCysProProCysLysCysProAla 279


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130 AAGGAGCGGTCTGGCGGCTTGAAGAAATTTGGACGATTTGGCAGCTTTGAGGCTCAAGGT 189
86 GlyLeuGlnAenIleAlaAaGluYshIshenLeuGlyIleLeuThrLysAraSerAen 105
190 GCATTGGCCAACTAGCTGTGGACAAAGCCAACTTGGAAATCATGACAAAGCGCTCCAAC 249
106 PheThrProAlaThrAsnGluAlaProGlnAlaThrValPheProLysSerProValLeu 125
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126 LeuGlyGlnProAenThrLeuIleCysPheValAspAenIlePheProProValIleAen 145
310 CTGAGAGCCCAACGCTCTCATCTGTGTTTCATAGACAAAGTTTCACCCACCAAGTGTGCAAT 369
146 IleThrTrpLeuArgAsnSerLysSerValThrAspGlyValThrGluThrSerPheLeu 165
370 GTCACGTGGCTTCGAAATGGAACAACTGTCAACAGAGGTGTGAGACAGATCTTCTCTG 429
166 ValAenArgAspHisSerPheHisLysLeuSerTyrLeuThrPheIleProSerAspAsp 185
430 CCCAGGGAAGACCACCTTTTCGCAAGTTCCACTATCTCCCTTCTCCCTCAACTGAG 489
186 AsplleTyrAspCysLysValGluHisTrpGlyLeuGluGluProValLeuLysHisTrp 205
490 GACGTTTACGACTGACGGGTGGAGCACTGGGGCTTGGATGAGCCTCTTCTCAAGCAGCTGG 549
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550 GAGTTTGATGTCCTCAAGCCCTCTCCAGAGACTACAGAGGTCGACGGAGGTGGCGCGGT 609
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610 TTAACGTATACACTCCAAGCGGAGACAGATCAACTTGAAGAGAGAGATCTGGTGGAG 669
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260 AlaAlaSerGluProArgGlyProThrIleLysProCysProCysLysCysProAla 279
730 GCAGCATCTGAGCCCAAGAGGCGCCCAATCAAGCCCTGTCTCCATGCAATGCGCCAGCA 789
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850 ATGATCTCCCTGAGCCCATAGTCATGATGTGGTGGTGGATGTGAGCGAGGATGACCCA 909
330 AspValGlnIleSerTrpPheValAenAenValGluValHisThrAlaGlnThrGlnThr 339
910 GATGTCCAGATCAGCTGGTGTGTGAAACAACTGGGAAGTACACACAGCTCAGACACAAACC 969
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1090 ATCGAGAGAACCATCTTCAAAACCCAAAGGTCAGTAAAGGCTCCACAGGTATATGTCTTG 1149
400 ProProProGluGluGluMetThrLysLysGlnValThrLeuThrCysMetValThrAsp 419
1150 CCTCCACCAAGAAGAGATGACTAAGAAACAGGTCACCTGACCTGATGCTGTCACAGAC 1209
420 PheMetProGluAspIleTyrValGluTrpThrAenAenGlyLysThrGluLeuAenTyr 439
1210 TTCATGCTGAAGACATTTACGTGGAGTGGACCAACAAACGGGAAACAGAGCTAAACTAC 1269
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Db 1270 AAGNACACTGACACAGCTCCCTGGACTCTGATGGTTCTTACTTTCATGTACAGCAAGCTGAGA 1329
Qy 460 ValGluLysLysAenTrpValGluArgAenSerTyrSerCysSerValValHisGluGly 479
Db 1330 GTGGAAAAGAAGAACTGGGTGGAAAAGAAATAGCTACTCTCTGTTTCAGTGGTCCACGAGGGT 1389
Qy 480 LeuHisAenHisThrThrLysSerPheSerArgThrProGlyLys 495
Db 1390 CTGCACAATCACACACAGCTTAAGAGCTTCTCCCGGACTCCGGGTAAA 1437
RESULT 5
ADM44282
ID ADM44282 standard; DNA; 1446 BP.
XX
AC ADM44282;
XX
DT 24-MAR-2005 (first entry)
XX
DE DR2-IgG fusion protein encoding DNA.
XX
KW Major histocompatibility complex; fusion protein; immunoconjugate;
KW adoptive immunotherapy; dermatological; immunosuppressive; anti-rheumatic;
KW anti-arthritis; neuroprotective; anti-inflammatory; autoimmune diseases;
KW pemphigus vulgaris; rheumatoid arthritis; multiple sclerosis;
KW systemic lupus erythematosus; immune disorder; DR2-IgG protein; gene; ds.
XX
OS Homo sapiens.
OS Chimeric.
OS Unidentified.
XX
FH Key Location/Qualifiers
CDS 1..1440
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FT /product= "DR2-IgG fusion protein"
FT /partial
FT /note= "No start codon"
FT misc_feature 1..15
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FT /*tag= d
FT /note= "Linker sequence"
FT misc_feature 610..729
FT /*tag= e
FT /note= "Fos Leucine zipper domain"
FT misc_feature 730..1437
FT /*tag= f
FT /note= "IgG domain"
XX
US2005003431-A1.
XX
XX 06-JAN-2005.
XX
XX 21-JUL-2004; 2004US-00895543.
XX
XX 16-AUG-1996; 96US-0024077P.
XX 15-AUG-1997; 97WO-US014503.
XX 19-FEB-1998; 98US-0075351P.
XX 12-FEB-1999; 99US-00248964.
XX
XX (WUCH/) WUCHERPFENNIG K W.
XX (STRO/) STROMINGER J L.
XX
XX Wucherpennig KW, Strominger JL;
XX
XX WPI; 2005-089945/10.
XX P-PSDB; ADM44283.
```


27-SEP-2001.

22-MAR-2001; 2001WO-US009616.

22-MAR-2000; 2000US-0191274P.

15-MAY-2000; 2000US-0204249P.

23-JAN-2001; 2001US-0264003P.

(CORI-) CORIXA CORP.

Carter D, Zhu S, Aximilli S, Wang A;

WPI; 2001-616371/71.

P-PSDB; ABB56457.

Multimeric complex for treating autoimmune diseases, comprises first and second single chain MHC class II molecules, each comprising alpha1 and beta1 domain linked through amino acid linker and multimerization domain.

Disclosure: Page 91-92; 147pp; English.

The invention relates to a multimeric complex comprising a first recombinant single chain major histocompatibility complex (MHC) class II molecule and a second recombinant single chain MHC class II molecule, each comprising an alpha domain and a beta domain linked through an amino acid linker and a multimerisation domain. The first and the second molecule are linked through the multimerisation domain to form a multimeric complex. The complex is useful for treating autoimmune diseases. It is useful for treating insulin dependent diabetes, multiple sclerosis, myasthenia gravis, pernicious anaemia, autoimmune encephalomyelitis (EAE), rheumatoid arthritis and systemic lupus erythematosus. The present invention encodes a single chain MHC class II molecule of the invention.

Sequence 2346 BP: 560 A; 663 C; 646 G; 477 T; 0 U; 0 Other;

Instrument Scores:

Document Scores:					
Id. No.:	1.77e-139	Length:	2346		
Size:	1738.00	Matches:	336		
Cent Similarity:	70.8%	Conservative:	44		
Local Similarity:	62.6%	Mismatches:	79		
Hy Match:	65.3%	Indels:	78		
	4	Gaps:	8		

: 0-048-116B-2 (1-495) x ABI99027 (1-2346)

22 GlyGly-----GluAspAspIleGluAlaAspHisValGlyPheTyrGlyThr 38
 766 GCGCGTTCTCGAGTGAAGACAGCATTTAGGCGCCGACGAGTGGGCGTCTATGGTACAAC 825
 39 ValTyrGlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeu 58
 826 GTATTATCAGTCTCTGGAGACATTTGCCAGTACACACATGATTTGATGGTGATGAGTGG 885
 59 PheTyrValAspLeuAspLysLysLysThrValTrpArgLeuProGluPheGlyGlnLeu 78
 886 TTCTATGTGGACTTGGATATAAGAGGAGACTATCTGGATGCTTCTCGAGTTTGGCCAAATTG 945
 79 IleLeuPheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGly 98
 946 ACAAGCTTTTGACCCCCCAAGGTGGAGTCTCAAAACATAGCTACAGGAAAATACACCTTTGGGA 1005
 99 IleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrVal 118
 1006 ATCTTGACTAAGAGGTCAAATTTCCACCCAGCTACCAATAGGCTCCTCAAGCGACTGTG 1065
 119 PheProLysSerProValLeuLeuGlyGlnProAsnThrIleuIleCysPheValAspAsn 138
 1066 TTCCCCCAAGTCCCCTGTGCTGTGGGTACGCCCAACACCCCTCATCTGCTTTGTGGACAAAC 1125
 139 IlePheProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGly 158

459 ArgValGluLysLysSerTrpValGluArgAsnSerTyrSerCysSerValValHisGlu 478
 2221 AATGTGCAAGAGCAACTGGGAGGCGAGAAATACATTTTCCCTGCTCTGTGTACATGAG 2280

479 GlyLeuHisAsnHisHisThrThrLysSerPheSerArgThrProGlyLys 495

2281 GGCCTGCACACCACTACTGAGAGAGCCTCTCCCACTCTCCTGTGTA 2331

SULT 7

I99033

ABI99033 standard; cDNA; 2343 BP.

ABI99033;

25-FEB-2002 (first entry)

MBP 90-101 CH1.H.CH2.CH3 coding sequence.

Mouse; MHC; major histocompatibility complex; MHC class II; multimer;
 single chain; immunosuppressive; antidiabetic; antiinflammatory;
 antianemic; antirheumatoid; antiarthritic; neuroprotective; vaccine;
 autoimmune disease; insulin dependent diabetes; multiple sclerosis;
 myasthenia gravis; pernicious anaemia; autoimmune encephalomyelitis;
 rheumatoid arthritis; systemic lupus erythematosus; ss.

Mus sp.

Synthetic.

WO200170245-A1.

27-SEP-2001..

22-MAR-2001; 2001WO-US009616.

22-MAR-2000; 2000US-0191274P.

15-MAY-2000; 2000US-0204249P.

23-JAN-2001; 2001US-0264003P.

(CORI-) CORIXA CORP.

Carter D, Zhu S, Arimilli S, Wang A;

WPI; 2001-616371/71.

P-PSDB; ABB56463.

Multimeric complex for treating autoimmune diseases, comprises first and second single chain MHC class II molecules, each comprising alpha1 and beta1 domain linked through amino acid linker and multimerization domain.

Disclosure; Page 96; 147pp; English.

The invention relates to a multimeric complex comprising a first recombinant single chain major histocompatibility complex (MHC) class II molecule and a second recombinant single chain MHC class II molecule, each comprising an alpha1 domain and a beta1 domain linked through an amino acid linker and a multimerization domain. The first and the second molecule are linked through the multimerization domain to form a multimeric complex. The complex is useful for treating autoimmune diseases. It is useful for treating insulin dependent diabetes, multiple sclerosis, myasthenia gravis, pernicious anaemia, autoimmune encephalomyelitis (EAE), rheumatoid arthritis and systemic lupus erythematosus. The present sequence encodes a single chain MHC class II molecule of the invention

Sequence 2343 BP; 562 A; 665 C; 635 G; 481 T; 0 U; 0 Other;

ignment Scores:

sd. No.:	1.95e-139	Length:	2343
ore:	1737.50	Matches:	336
cent Similarity:	70.6%	Conservative:	44
at Local Similarity:	62.5%	Mismatches:	79
ary Match:	65.3%	Indels:	79

DB:	4	Gaps:	8
US-10-048-116B-2 (1-495) x ABI99033 (1-2343)			
Qy	22 GlyGly-----GluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrThr 38		
Db	760 GCGCGTTCCTCGAGTGAAGACGACATTTAGGCGCGACACGCTAGGCGCTCTATGTACAACCT 819		
Qy	39 ValTyrGlnSerProGlnGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeu 58		
Db	820 GTATATCAGTCTCCTGGAGACATTTGCCAGTAGTACACATGAATTTTATGGTGTATGATGG 879		
Qy	59 PheTyrValAspLeuAspLysLysThrValTyrArgLeuProGluPheGlyGlnLeu 78		
Db	880 TTCTATGTGACTTGGATAAGAGAGACTATCTGGATGCTTCTGAGTCTTGGCCAAATTG 939		
Qy	79 IleLeuPheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisLeuGly 98		
Db	940 ACAAGCTTTGACCCCAAGTGGACTGCAAAACATAGCTACAGGAAATATACACCTTTGGGA 999		
Qy	99 IleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrVal 118		
Db	1000 ATCTTGACTTAAGAGGTCAAAATTTCCACCCAGCTACCAATGAGGCTCTCAAGCGACTGG 1059		
Qy	119 PheProLysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsn 138		
Db	1060 TTCCCAAGTCCCTGTGCTGTGGTTCAGCCCAACACCTCTCATCTGCTTTGTGGACAC 1119		
Qy	139 IlePheProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGly 158		
Db	1120 ATCTTCCCTCTGTGATCAACATCATGCTGCTCAGAAATAGTAAGTCAGTCACAGACGGC 1179		
Qy	159 ValTyrGluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeu 178		
Db	1180 GTTATGAGACAGCTTCTGTCAACCGTGACCATCTCTCCACAGCTGCTTATCTC 1239		
Qy	179 ThrPheIleProSerAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGlu 198		
Db	1240 ACCTTCATCCCTTCTGACATGATATTTATGACTGCAAGTGGAGCAGCTGGGCGCTGGAG 1299		
Qy	199 GluProValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGlu 218		
Db	1300 GAGCGGTTCTGAACACACTGG-----GCT 1323		
Qy	219 ThrGlyGlyGlyGlySer-----ThrThrAlaProSer----- 229		
Db	1324 AGCGAGGGGGCGAAGCGCGGAGAGAGCTTAGCCAAACAGCACACCCCATCTGTCTAT 1383		
Qy	229 ----- 229		
Db	1384 CCACCTGGCCCCCTGGATCTGCTGCCCAACTAACTCCATCGTGTGACCTGGGATCGCTGGTC 1443		
Qy	229 ----- 229		
Db	1444 AAGGGCTATTTCCCTGAGCCAGTGACAGTGACCTTGGAACTCTGGATCCCTGTCCAGCGGT 1503		
Qy	230 -----AlaGlnLeuGluLysGluLeuGlnAlaLeuGluLysGluAsnAla 244		
Db	1504 GTGCACACCTTCCAGCTGCTCTGCACTGTGACCTCTACACCTTGAGCAGCTCAGTGACT 1563		
Qy	245 -----GlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeuAlaGlnAlaLaser 262		
Db	1564 GTCCCTCCAGCACCTGGCCCCAGGAGACCGTACCTGCAACGTTGCCACCCCGGCAGC 1623		
Qy	263 GluProArg---GlyProThrIleLysPro-----CysProProCysLysCys 277		
Db	1624 AGCACCAAGGTGGACAAGAAATTTGGCCAGGAGTTGGTTGTGAAGCTTGCATATGT 1683		
Qy	278 ProAlaProAsnLeuLeuGlyGlyProSerValPheIlePheProProLysIleLysAsp 297		
Db	1684 ACAGTCCCAAGAAGTA-----TCATCTGTCTTTCATCTTCCCCCAAGCCCAAGGAT 1734		
Qy	298 ValLeuMetIleSerLeuSerProIleValThrCysValValValAspValSerGluAsp 317		

219 ThrGlyGlyGlySer-----ThrAlaProSer----- 229
 1324 AGCGAGGGGCGAAGCGGCGGAGGAGCAAAACAAACACACCCCTCATCTATCCA 1383
 229 ----- 229
 1384 CTGGCCCTGGGTGGAGATACAACCTGGTCTCGGTGACTCTGGGATGCCTGTCAG 1443
 229 ----- 229
 1444 GGTACTTCCCTGAGTCACTGAGTCTGGAACTCTGGCTCCCTGTCCAGCAGTGTG 1503
 230 -----AlaGlnLeuGluysGluLeuGlnAlaLeuGluysGluAla--- 244
 1504 CACACCTTCCCAAGCTCTCTCGAGTCTGGACTCTACATATGAGCAGCTCACTGTGTC 1563
 245 ---GlnLeuGluTTPGluLeuGlnAlaLeuGluysGluLeuAlaAlaSer--- 262
 1564 CCTTCAGCAGCTGGCAAGTCAGACCTGACCTGACCTGAGGGTGTCTACCCAGCAGCAGC 1623
 263 -----GluProArgGlyPro-----ThrIleLysProCysPro 273
 1624 ACCACGGTGGCAAAAACCTTGAGCCAGCGGCCCATTTCAACAATCAACCCCTGTCT 1683
 274 ProCys-----LysCysProAlaProAsnLeuGluGlyGlyProSerValPhe 289
 1684 CCATGCAAGGAGTGTACAAAATGCCAGCTCTTAACCTGGAGGGTGGACCATCCGTCTTC 1743
 290 IlePheProLysIleLysAspValLeuMetIleSerLeuSerProIleValThrCys 309
 1744 ATCTTCCCTCCAAATATCAAGATGATCATGATCTCTCTGACACCCAGGTCAAGTGT 1803
 310 ValValValAspValSerGluAspProAspValGlnIleSerTrpPheValAsnAsn 329
 1804 GTGGTGGTGGATGTGCGAGGAGTACCCAGAGCTCCAGATCAGTGTGTTGTGAACAAC 1863
 330 ValGluValHisThrAlaGlnThrGlnThrHisArgGluAspTrpAsnSerThrLeuArg 349
 1864 GTGGAAGTACACACAGCTCAGACACAAACCCATAGAGAGGATTACACAGTACTATCCGG 1923
 350 ValValSerAlaLeuProIleGlnHisGlnAspTrpMetSerGlyLysGluPheLysCys 369
 1924 GTGGTCAGACCCCTCCCATCCAGACCCAGGACTGGATGAGTGGCAGAGGTTCANATGC 1983
 370 LysValAsnAsnLysAspLeuProAlaProIleGluArgThrIleSerLysProLys 388
 1984 AAGGTCAACAAAGACCTCCCATCCATCCAGAGAACCATCTCAAAAATTTAA 2040

SULT 9
 199032

ABI99032 standard; cDNA; 2059 BP.

ABI99032;

25-FEB-2002 (first entry)

MBP 1-14 Ch1.H.CH2 coding sequence.

Mouse; MHC; major histocompatibility complex; MHC class II; multimer; single chain; immunosuppressive; antidiabetic; antiinflammatory; antianemic; antirheumatoid; antiarthritic; neuroprotective; vaccine; autoimmune disease; insulin dependent diabetes; multiple sclerosis; myasthenia gravis; pernicious anaemia; autoimmune encephalomyelitis; rheumatoid arthritis; systemic lupus erythematosus; ss.

Mus sp.
 Synthetic.

WO200170245-A1.

27-SEP-2001.

PF 22-MAR-2001; 2001WO-US009616.
 XX 22-MAR-2000; 2000US-0191274P.
 PR 15-MAY-2000; 2000US-0204249P.
 PR 23-JAN-2001; 2001US-0264003P.
 XX (CORI-) CORIXA CORP.
 XX Carter D, Zhu S, Arimilli S, Wang A;
 PI WPI; 2001-616371/71.
 DR P-PSDB; ABB56462.
 XX Multimeric complex for treating autoimmune diseases, comprises first and second single chain MHC class II molecules, each comprising alpha and beta domain linked through amino acid linker and multimerization domain.
 PT Disclosure; Page 95; 147pp; English.
 XX The invention relates to a multimeric complex comprising a first recombinant single chain major histocompatibility complex (MHC) class II molecule and a second recombinant single chain MHC class II molecule, each comprising an alpha domain and a beta domain linked through an amino acid linker and a multimerization domain. The first and the second molecule are linked through the multimerization domain to form a multimeric complex. The complex is useful for treating autoimmune diseases. It is useful for treating insulin dependent diabetes, multiple sclerosis, myasthenia gravis, pernicious anaemia, autoimmune encephalomyelitis (RAE), rheumatoid arthritis and systemic lupus erythematosus. The present sequence encodes a single chain MHC class II molecule of the invention
 XX SQ Sequence 2059 BP; 493 A; 585 C; 571 G; 410 T; 0 U; 0 Other;
 Alignment Scores:
 Pred. No.: 7,22e-121 Length: 2059
 Score: 1521.00 Matches: 304
 Percent Similarity: 72.0% Conservative: 12
 Best Local Similarity: 69.2% Mismatches: 39
 Query Match: 57.2% Indels: 84
 DB: 4 Gaps: 8
 US-10-048-116B-2 (1-495) x ABI99032 (1-2059)
 QY 22 GlyGly-----GluAspAspIleGluAlaAspHisValGlyPheTrpGlyThrThr 38
 Db 766 GCGCGTTCCTCGAGTGAAGACGACATTGAGGCCGACCGTAGGCGTCTATGGTACAACT 825
 QY 39 ValTrpGlnSerProGlyAspIleGlyGlnTrpThrHisGluPheAspGlyAspGluLeu 58
 Db 826 GTATATCAGTCTCTCGGAGACATTGGCCAGTACACACATGAATTTGATGGTGTAGTGG 885
 QY 59 PheTrpValAspLeuAspLysLysLysThrValTrpArgLeuProGluPheGlyGlnLeu 78
 Db 886 TTCTATGTGGACTTGGATAAGAAGGAGACTATCTGGATGTTCTCTGAGTTGGCCAAATG 945
 QY 79 IleLeuPheGluProGlnGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGly 98
 Db 946 ACAAGCTTGGACCCCAAGGTGGACTGCAAAACATAGCTACAGGAAAATACACCTTTGGGA 1005
 QY 99 IleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrVal 118
 Db 1006 ATCTTGACTAGAGGTCAAAATTCACCCAGTACCAATAGGCTCTCAAGGACTGTG 1065
 QY 119 PheProLysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsn 138
 Db 1066 TTCCCAAGTCCCTGTGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 1125
 QY 139 IlePheProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGly 158
 Db 1126 ATCTTCCCTCTGTGTATCAACATCATGTGGCTCAGAAAATAGTAAAGTCAAGACGGC 1185
 QY 159 ValTrpGluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTrpLeu 178

1186 GTTTATGAGACCACTTCTTGTTCACCGGTGACCATTTCTTCCACAGCTGTCTTATCTC 1245
179 ThrPhe11eProSerAspAspAsp1e1yAspCysLysValGluHisTrpGlyLeuGlu 198
1246 ACCTTCATCCCTTCTGACGATGATATTTATGACTGCAAGGTGGAGCACTGGGGCTGGAG 1305
199 GluProValLeuLysHisTrpGluProGluLeuProAlaProMetSerGluLeuThrGlu 218
1306 GAGCCGGTCTCTGAACAACATGG-----GCT 1329
219 ThrGlyGlyGlyGlySer-----ThrThrAlaProSer----- 229
1330 AGCGAGGGGGCGAGAGCGGGAGGGAGGCCAAACACACACCCCATCAGTCTATCCA 1389
229 ----- 229
1390 CTGGCCCTTGGGTGTGGAGATACAACCTGGTTCTCTCCGTGACTCTGGGATGCTGCTCAAG 1449
229 ----- 229
1450 GGCTACTTCCCTGAGTCAGTGACCTGTGACTTTGGAACTCTGGCTCCCTGTCGACGAGTGTG 1509
230 -----AlaGlnLeuGluLysGluLeuGlnAlaLeuGluLysGluAsnAla--- 244
1510 CACACCTTCCAGCTCTCTCTGACTCTGGACTCTACACTATGAGCAGCTCAGTGACTGTC 1569
245 ---GlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeuAlaGlnAlaSer--- 262
1570 CCCTCCAGCACCTTGGCCAAAGTCAGACCGTCACTCTGCAGCGTTGTCTACCCAGCCAGCAGC 1629
263 -----GluProArgGlyPro-----ThrLeuLysProCysPro 273
1630 ACCAGGTGGCAAAAAAATTTGAGCCAGCGGGCCATTTCAACAATCAACCCCTGTCTCT 1689
274 ProCys-----LysCysProAlaProAsnLeuLeuGlyGlyProSerValPhe 289
1690 CCATGCAAGGAGTGTCAAAATGCCAGCTCTAAACCTGGAGGGTGAGCACCATCCGCTCTC 1749
290 IlePheProProLysIleLysAspValLeuMetIleSerLeuSerProIleValThrCys 309
1750 ATCTTCCCTCCAAATATCAAGATGTAATCATGATCTCCCTGACCAACCAAGTCACTGTGT 1809
310 ValValValAspValSerGluAspAspProAspValGlnIleSerTrpPheValAsnAsn 329
1810 GTGTGTGTGATGTGTGAGCGAGGATGATCCAGACGCTCCAGATCAGTCTGTTGTGCAACAC 1869
330 ValGluValHisThrAlaGlnThrGlnThrHisArgGluAspTyrAsnSerThrLeuArg 349
1870 GTGGAAGTACACACAGCTTCAGACACAACCCATAGAGAGGATTAACAACAGTACTATCCGG 1929
350 ValValSerAlaLeuProIleGlnHisGlnAspTrpMetSerGlyLysGluPheLysCys 369
1930 GTGGTCAGCACCTTCCCATCCAGCACGAGGATGGATGATGGCAGAGGAGTTCAAATGC 1989
370 LysValAsnAsnLysAspLeuProAlaProIleGluArgThrIleSerLysProLys 388
1990 AAGGTCAACAACAAGAAGCCTCCCATCAACCCATCGAGAGAACCATCTCAAAAATTTAA 2046

ULT 10
:0316
AAF30316 standard; cDNA; 7528 BP.
AAF30316;
11-SEP-2003 (revised)
14-MAY-2001 (first entry)
Bicistronic idiotype plasmid VR1642.
Flt-3 ligand; Fms-like tyrosine kinase; mouse; human; vaccine;
immunotherapy; therapy; tumour; lymphoma; gene therapy; VR1642;
plasmid VAX1D; antibody; idiotype; vector; ss.

88 GlnAsnIleAlaLeuGlyHisLeuThrLysArgSerAsnPheThr 107
1613 GACGGG-----AGGTACCGTATGGACGCTTGGGGCCAAAGGACCAAGCTCAGC 1660
108 ProAlaThrAsnGluAlaProGlnAlaThrValPheProLysSerProVal 124
1661 GTCTCTCGGCCAAACGGCCGCCCTATCCCTATCCCTGCTGCTGGAGAT 1720
125 LeuLeuGlyGlnProAsnThrLeuLeuCysPheValAspAsnIlePheProValIle 144
1721 ACAAGTGGCTCTCGGTGCTAGTATGCTGGTCAAGGGTTATTCCTGAGCCAGTG 1780
145 AsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValThrSerPhe 164
1781 ACCTTGACCTGG-----AACTCTGGATCCCTGCTCAGTGGTGCACACCTTCCAGCT 1834
165 LeuValAsnArgAspHisSerPheHisLysLeuSerThrLeuThrPheIleProSerAsp 184
1835 GTCTGTCAGCTGAC-----CTCTACACCTCAGCAG-CTCAGT----- 1872
185 AspAspIleTyrAspCysLysValGluHisTrpGlyLeuGluGluProValLeuLysHis 204
1873 -----GACTGTAACTCGAGCAC-----CTGGCCAGCAGTC-----CAT 1908
205 TrpGluProGluIleProAlaProMetSerGluLeuThrGluThrGlyGlyGly-Se 224
1909 TAC-----CTGCAATGTGGGCCACCCGGCAAG 1935
224 rThrAlaProSerAlaGlnLeuGluLysGluLeuGlnAlaLeuGluLysGluAsnAl 244
1936 CAGCACC-----AAGGTGGACAGAAATTT----- 1960
244 aGlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeuAlaGlnAlaSerGluPr 264
1961 -----GAGCC 1965
264 ohArgGlyProThrIleLysProCysProCysLysCysProAlaProAsnLeuGlu 284
1966 CAGAGGGGCCCAACATCAAGCCCTGCTCCATGCAAAATGCCAGCACCTTAACCTTGGG 2025
284 yGlyProSerValPheIlePheProProLysIleLysAspValLeuMetIleSerLeuSe 304
2026 TGGACCATCCGCTTCCTCACTTCCTCCAAAGATCAAGGATGCTACTGATCTCCCTGAG 2085
304 rProIleValThrCysValValAspValSerGluAspAspProAspValGlnIleSe 324
2086 CCCCATAGTCACATGTGTGGTGGATGTGAGCGAGGATGACCCAGATGTCAGATCAG 2145
324 rTrpPheValAsnAsnValGluValHisThrAlaGlnThrGlnThrHisArgGluAspTy 344
2146 CTGGTTTGTGAACAACGTGGAGATACACACAGCTCAGACACAAACCCATAGAGAGGATTA 2205
344 rAsnSerThrLeuArgValValSerAlaLeuProIleGlnHisGlnAspTrpMetSerG1 364
2206 CAACAGTACTCTCCGGTGGTCAAGTGGCTCCCTCCATCCAGCACAGGACTGGATGATGG 2265
364 yLysGluPheLysCysLysValAsnAsnLysAspLeuProAlaProIleGluArgThrI1 384
2266 CAAGGAGTTCAATGCAAGGTCAACAACAAGACCTCCAGCGCCCATCAGAGAAGAACAT 2325
384 eSerLysProLysGlySerValArgAlaProGlnValThrValLeuProProGluG1 404
2326 CTCAAACCCCAAGGGTCAAGAGCTCCACAGGTATATGCTTGGCTCCACCAAGAAGA 2385
404 uGluMetThrLysLysGlnValThrLeuThrCysMetValThrAspPheMetProGluAs 424
2386 AGAGATGACTAAGAAACAGGTCACTCTGACCTGCAATGGTCAACAGACTTCATCCCTGAAGA 2445
424 pIleTyrValGluTrpThrAsnAsnGlyLysThrGluLeuAsnTyrLysAsnThrGluPr 444
2446 CATTTACGTGGAGTGGACCAACAACGGGAAACAGAGCTAAACTCAAGAACACTGAACC 2505
444 oValLeuAspSerAspGlySerTyrPheMetTyrSerLysLeuArgValGluLysLysAs 464

Db 2506 AGTCTGAGCTCTGATGTTCTTACTTATGTACAGCAAGCTGAGAGTGGAAAGAGAA 2565
Qy 464 nTrpValGluArgAsnSerTyrSerCysSerValValHisGluGlyLeuHisAsnHisH1 484
Db 2566 CTGGGTGAAAGAAATAGTACTACTCTCTTCTAGTGTCCACGAGGGTCTGCACATCACCA 2625
Qy 484 sThrThrLysSerPheSerArgThrProGlyLys 495
Db 2626 CACGACTAAGAGCTTCTCCGAGCTCCGGTAA 2659
RESULT 11
ADT77690
ID ADT77690 standard; DNA; 3973 BP.
XX
AC ADT77690;
XX
DT 13-JAN-2005 (first entry)
XX
DE Monoclonal antibody mAb17-1A expression construct.
XX
KW Mouse; IgG2a; antibody; monoclonal antibody; mAb17-1A; cancer; vaccine;
KW gene; ds.
XX
OS Mus sp.
XX
FH Key Location/Qualifiers
FT CDS 25..756
FT /*tag= a
FT /product= "mAb17-1A kappa chain"
FT 1357..2754
FT /*tag= b
FT /product= "mAb17-1A gamma chain"
XX
PN WO2004091655-A2.
XX
PD 28-OCT-2004.
XX
PF 16-APR-2004; 2004WO-EP004059.
XX
PR 17-APR-2003; 2003AT-00000599.
XX
PA (IGEN-) IGENEON KREBS IMMUNOTHERAPIE FORSCHUNGS.
XX
PI Loibner H, Himmler G, Waxenecker G, Schuster M, Putz T;
XX
DR WPI; 2004-758278/74.
DR P-PSDB; ADT77691, ADT77692.
XX
DR New immunogenic recombinant antibody comprising a part of a murine IgG2a
PT subtype amino acid sequence and a mammalian glycosylation, useful in
PT preparing a vaccine for immunizing primates against infectious or
PT autoimmune disorders.
XX
PS Disclosure; SEQ ID NO 1; 59pp; English.
XX
CC The present sequence is that of the cloning cassette of a tri-clonronic
CC mAb17-1A expression and dihydrofolate reductase selection construct for
CC use in the production of recombinant IgG2a mAb17-1A antibody. In examples
CC from the invention, comparison of the original, hybridoma-derived
CC immunisation antigen 17-1A and recombinantly expressed mAb17-1A from CHO
CC cells did not reveal any immunological differences in rhesus monkey
CC trials. Both formulations showed identical kinetics building up the
CC immunisation antigen and target antigen specific immune response, and
CC raised IgG and IgM titres were similar. Recombinant mAb17-1A is an
CC example of an immunogenic recombinant antibody of the invention designed
CC for immunisation of primates and comprising at least a part of a murine
CC IgG2a subtype amino acid sequence with mammalian glycosylation.
CC Recombinant DNA methods are used to produce the immunogenic antibody in a
CC standardised manner. The humoral immune response induced by the IgG2a
CC immunogenic antibodies of the invention is significantly improved in
CC terms of the quantity of specific antibody induced by the patients and
CC the specificity against selected targets and epitopes. The improved

immune response is dependent on the glycosylation pattern of the antibody. Recombinant antibody expressed in hamster or human cells shown to have a similar immunogenicity as antibody expressed by murine hybridoma cells. This is of particular relevance for antibodies that are to be used for immunisation purposes. The antibody may have a murine amino acid sequence or any other mammalian amino acid sequence that is combined with the murine IgG2a part. Preferably mammalian sequences are human, humanized, human/murine chimeric or murine sequences. The antibody may also be an anti-idiotypic antibody or a mimotopic Abi antibody. The IgG2a immunogenic recombinant antibody can be directed against a tumour associated antigen. The invention also provides vaccines comprising the IgG2a immunogenic antibody. The vaccines may be used for the prophylaxis and therapy of cancer associated diseases, e.g. metastatic disease in cancer patients. The vaccine specifically modulates antigen-presenting cells in vivo or ex vivo, thus generating an immune response to the epitope that is targeted by the IgG2a immunogenic antibody. The preferred method of producing the antibody comprises: transforming a CHO host cell with a multicistronic antibody-expression construct containing at least a nucleotide sequence encoding a kappa light chain and a nucleotide sequence encoding a gamma heavy chain, where at least one of these sequences comprises a nucleotide sequence encoding at least a part of a murine IgG2a subtype amino acid sequence, and at least 2 IRES elements; and expressing the sequences under the control of a single CMV promoter to produce an intact antibody. The kappa light chain and gamma heavy chain are expressed in about equimolar quantity, and antibody concentrations of 5-300 ug/ml are achieved.

Sequence 3973 BP; 1052 A; 1038 C; 993 G; 888 T; 0 U; 2 Other;

Alignment Scores:

d. No.:	6.48e-104	Length:	3973
Percent Similarity:	1328.50	Matches:	281
Local Similarity:	65.0%	Conservative:	26
Match:	59.5%	Mismatches:	71
	49.9%	Indels:	95
	13	Gaps:	11

: 0-048-116B-2 (1-495) x ADT77690 (1-3973)

```

42 SerProGlyAspIleGly---GlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
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1567 AATCTGGAAAGTGGTGACTACTACAAATGAGAAGTTCAAGGGCAAGGCAACACTGACT 1626
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
61 ValAspLeuAspLysLysLysThrValTyrArgLeuProGluPhe----- 75
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1627 GCAGACAAATCTCCAGCACTGCCTACATGAGCTCAGCTCAGCAGCTGATGACTCT 1686
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
76 -----GlyGlnLeuIleLeuPheGluProGlnGlyGlyLeu 87
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1687 GCGGTCTATTCTGTGCAAGAGATGGTCCCTGGTTTGTCTTACTGGGGCCAGGGACTCTG 1746
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
88 GlnAsnIleAlaAlaGluLysHisAsnLeuGlyIleLeuThrLysArgSerAsnPheThr 107
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1747 GTCACTGTCTCTGCAGCCCAA----- 1767
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
108 ProAlaThrAsnGluAlaProGlnAlaThrValPheProLysSerProVal----- 124
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1768 -----ACACAGCCCA-----TCGGTCTATCCACTGGCCCTGTGTGGAGAT 1812
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
125 LeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePheProValIle 144
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1813 ACAACTGGCTCTCGGTGACTTAGGATGCTGGTCAAGGGTTATTTCCCTGAGCCAGTG 1872
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
145 AsnIleThrTropLeuArgAsnSerLysSerValThrAspGlyValTyrGluThrSerPhe 164
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1873 ACCTTGACCTGG-----AATCTGGATCCCTGCTCAGTGGTGACACCTTCCCGACT 1926
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
165 LeuValAsnArgAspHisSerPheHisLysLysSerTyrLeuThrPheIleProSerAsp 184
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1927 GTCTGCACTGAC-----CTACACCTCAGCAG-CTCAGT----- 1964
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
185 AspAspIleTyrAspCysLysValGluHis-TripGlyLeuGluProValLeuLysHis 204
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

```

Db	1965	-----GACTGTAACTCGAGCACCTGG-----	1986
Qy	204	sTrpGluProGluIleProAlaProMetSerGluLeuThrGluThrGlyGlyGlyGlyse	224
Db	1987	-----CCAGCCAGTCCATCACC-----	2015
Qy	224	rThrAlaProSerAlaGlnLeuGluLysGluLeuGlnAlaLeuGluLysGluAsnAl	244
Db	2016	CCACCCGCAAGCAGCACCACCAAGGTGGCAAGAAATT-----	2052
Qy	244	aGlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeuAlaAlaSerGluPr	264
Db	2053	-----GAGCC-----	2057
Qy	264	oAtsGlyProThrIleLysProCysProCysLysCysProAlaProAsnLeuLgl	284
Db	2058	CAGAGGCCCAATCAATCAAGCCCTGCTCCATGCAATGCCAGCACCTAACCTCTTGGG	2117
Qy	284	yGlyProSerValPheIlePheProProLysIleLysAspValLeuMetIleSerLeuSe	304
Db	2118	TGGACCATCGGTCTTCACTTCCCTCCAAAGATCAAGGATGTACTCATGTCTCCCTGAG	2177
Qy	304	rProIleValThrCysValValValAspValSerGluAspAspProAspValGlnIleSe	324
Db	2178	CCCATAGTCACATGTGTGGTGGATGTGAGCGAGATGACCCAGATGTCCAGATCAG	2237
Qy	324	rTrpPheValAsnAsnValGluValHisThrAlaGlnThrGlnThrHisArgGluAspTy	344
Db	2238	CTGGTTGTGAACACCTGGAGTACACACAGCTCAGACACAAACCCATAGAGAGGATTA	2297
Qy	344	rAsnSerThrLeuArgValValSerAlaLeuProIleGlnHisGlnAspTrpMetSerGl	364
Db	2298	CAACAGTACTCTCGGGTGTGAGTGGCCCTCCCATCCAGCACCCAGGACTGGATGGTG	2357
Qy	364	yLysGluPheLysCysLysValAsnAsnLysAspLeuProAlaProIleGluArgThrIl	384
Db	2358	CAAGGAGTTCAATGCAAGGTCAACAAAGAGCTCCAGCGCCCATCGAGAGAACCAT	2417
Qy	384	eSerLysProLysGlySerValArgAlaProGlnValTyrValLeuProProGluGl	404
Db	2418	CTCAAAACCCAAAGGTGAGTCCAGAGTATATGTCTTGGCTCCACCAAGAGA	2477
Qy	404	uGluMetThrLysLysGlnValThrLeuThrCysMetValThrAspPheMetProGluAs	424
Db	2478	AGAGATGACTAAGAAAACAGGTCTACTGTGACTGTGATGGTCCACAGACTTCATGCTGAAGA	2537
Qy	424	pIleTyrValGluTrpThrAsnAsnGlyLysThrGluLeuAsnTyrLysAsnThrGluPr	444
Db	2538	CAITTAGTGGAGTGGACCAACACGGGAAACAGAGCTAAACTCAAGAACACTGAAC	2597
Qy	444	oValLeuAspSerAspGlySerTyrPheMetTyrSerLysLeuArgValGluLysLysAs	464
Db	2598	AGTCTGAGCTCTGATGGTTCTTACTTTCATGTACAGCAAGCTGAGAGTGGAAAGAGAA	2657
Qy	464	nTrpValGluArgAsnSerTyrSerCysSerValHisGluGlyLeuHisAsnHisHi	484
Db	2658	CTGGGTGGAAAGAAATAGTACTCTCTGTTCAGTGGTCCACAGGGGTCTGCACAATCACA	2717
Qy	484	sThrThrLysSerPheSerArgThrProGlyLys	495
Db	2718	CACGACTAAGAGCTTCTCCCGGACTCCCGGGTAAA	2751

RESULT 12

AAQ48037
ID AAQ48037 standard; cDNA to mRNA; 1581 BP.

AC AAQ48037;

XX 25-MAR-2003 (revised)

DT 10-MAR-2003 (revised)

DT 08-FEB-1994 (first entry)

DE Monoclonal antibody M(alpha)2-3 H-chain coding sequence.

anti-snake small neurotoxin antibody; heavy chain; IgG2; immunoglobulin; bispecific bivalent antibody; cell-targeting; cytotoxic agent; ss.

Unidentified.

Key	Location/Qualifiers
CDS	61..1470
	/*tag= b
sig_peptide	/product= "Ig_heavy-chain"
	61..117
mat_peptide	/*tag= a
	118..477
	/*tag= c
mat_peptide	/product= "Ig_variable_region"
	478..768
	/*tag= d
mat_peptide	/product= "Ig_constant_region"
	769..816
	/*tag= e
mat_peptide	/product= "Ig_joining_region"
	817..1146
	/*tag= f
mat_peptide	/product= "Ig_constant_region"
	1147..1167
	/*tag= g
	/product= "Ig_constant_region"

EP556111-A1.

18-AUG-1993.

09-FEB-1993: 93EP-00400323.

11-FEB-1992: 92FR-00001505.

(COMS) COMMISSARIAT ENERGIE ATOMIQUE.

Boulain J, Ducancel F, Gillet D, Menez A:

WPI; 1993-260351/33.
P-PSDB: AAR40384.

New immunoglobulin hybrid proteins - with immunoglobulin fragments linked to dimeric protein, for diagnostic or therapeutic use.

Example 1; Fig 3A; 37pp; French.

A fragment of the heavy chain (VH + CH1) from the anti-snake small neurotoxin monoclonal antibody M(alpha)2-3 was PCR-amplified from hydromedusa-derived cDNA using primers AAQ48039 and AAQ48040. A light chain fragment (VL + CL) was amplified from the same source using primers AAQ48041 and AAQ48042. The two amplified fragments were inserted into the same vector; the H-chain fragment was inserted (in-frame) between codons 6-7 of the phoA coding sequence and the L-chain fragment was inserted into a cassette which contained a phoA S-D sequence, a signal peptide and the first 6 codons of phoA. The cassette was positioned between the termination codon and the transcription termination sequence of phoA. The fusion construct is expected to encode a hybrid protein comprising two identical Ab-derived units. The invention also covers hybrid proteins containing two different Ab-derived units (i.e. to produce bispecific antibodies). When a toxic protein is used in place of phoA, the hybrid molecules can be used as cell-targeting therapeutic agents. (Updated on 10-MAR-2003 to add missing OS field.) (Updated on 25-MAR-2003 to correct PN field.)

Sequence 1581 BP: 435 A: 448 C: 373 G: 325 T: 0 U: 0 Other:

Assignment Scores:

Alignment scores:			
ed. No.:	4.24e-104	Length:	1581
ore:	1324.50	Matches:	286
ercent Similarity:	65.4%	Conservative:	31
st Local Similarity:	59.0%	Mismatches:	81

Query Match:	49.8%	Indels:	89
DB:	2	Gaps:	11
US-10-048-116B-2 (1-495) x AAQ48037 (1-1581)			
Qy	34	pheTyRGlyThrThrValTyRGlnSerProGlyAsp-----IleGly-----	47
Db	211	TACTATATAAACATGGGTGAAGCAAGACCTGGACAGGACTTAAATGGATGGATGGATT	270
Qy	48	-----GlnTyrThrHisGluPheAspGlyAspGluLeuPheTyR	60
Db	271	TATCCTGCAAGCGGTAACTAAGTAACTAGAACTTCAAGGCAAGGCCACATTGGACT	330
Qy	61	ValAspLeuAspIysLysThrValTrpArgLeu-----ProGluPhe	75
Db	331	GTAGACACATCTCCAGCACAGCTACATGCAGCTCAGCAGCCTGCATCTGAGGACACT	390
Qy	76	GlyGlnLeuIleLeuPheGluProGlnGlyGlyLeuGlnAsnIlealaGluIysHis	95
Db	391	GCTGT-CTATTTCTGCAAGAGCTATGGGGCTAC-----GGCTAC	431
Qy	96	AsnLeuGlyIleLeuThrLysArg-SerAsnPheThrProAlaThrAsnGluAlaProGln	115
Db	432	ACTTTTGGACTACTGGGCCNAGGCACCACTCTCACAGTCTCTTCAGCCAAACACACAGC	491
Qy	115	nAlaThrValPheProLysSerProVal-----LeuLeuGlyGlnProAsnThrIle	132
Db	492	CCCATCGGTCTATCCACTGGCCCTGTGTGTGAGATACAACTGGTCTCTCGGTGACTCT	551
Qy	132	uIleCysPheValAspAsnIlePheProProValIleAsnIleThrTrpLeuArgAense	152
Db	552	AGGATCCCTGGTCAAGGGTTATTTCCCTGAGCCAGTGACTTGGACTCG-----AACTC	605
Qy	152	ryLysSerValThrAspGlyValTyRGlnThrSerPheLeuValAsnArgAspHisSerPh	172
Db	606	TGGATCCCTGTCAGTGGTGTGCACACTCCAGCTGCTCCTGCAGTCTGAC-----CT	659
Qy	172	eHisLysLeuSerTyLeuThrPheIleProSerAspAspAspIleTyAspCysLysVa	192
Db	660	CTACACCCCTCAGCAG-CTCAGT-----GACTGTAACTCT	691
Qy	192	lGluHis--TrpGlyLeuGluGluProValLeuLysHisTrpGluProGluIleProAlap	212
Db	692	CGAGCACTGG-----C	703
Qy	212	roMetSerGluLeuThrGluThrGlyGlySerThrThrAlaproSerAlaGlnL	232
Db	704	CCAGCCAGTCCATCAC-----TGCAATGTGCCCCAGCCGCGCAGCAGCACCAAGG	754
Qy	232	euGluLysGluLeuGlnAlaLeuGluLysGluAsnAlaGlnLeuGluTrpGluLeuGlnA	252
Db	755	TGGACAAGAAAT-----	768
Qy	252	laLeuGluLysGluLeuAlaGlnAlaAlaSerGluProArgGlyProThrIleLysProC	272
Db	769	-----GAGCCCAAGGGCCCAATCAAGCCCT	796
Qy	272	ysProProCysLysCysProAlaProAsnLeuGlyGlyProSerValPheIlePheP	292
Db	797	GTCTCCATGCAATGCCAGCACCTAACTCTTGGGTGGACCATCCGCTCTTCATCTTCC	856
Qy	292	roProLysIleLysAspValLeuMetIleSerLeuSerProIleValThrCysValValV	312
Db	857	CTCCAAGATCAAGGATGTACTCATGATCTCCCTGAGCCCCATAGTCACATGTGTGGTGG	916
Qy	312	aAspValSerGluAspAspProAspValGlnIleSerTrpPheValAsnAsnValGluV	332
Db	917	TGGATGTGACGGAGGATGACCCAGATGTCAGATCAGCTGGTTTGTGAACACAGTGGAG	976
Qy	332	alHisThrAlaGlnThrGlnThrHisArgGluAspTyRAsnSerThrLeuArgValValS	352
Db	977	TACACACAGCTCAGACACAAACCAATAGAGAGATTACACAGTACTCTCCGGGTGGTCA	1036

352 eAlaLeuProIleGlnHisGlnAspTyrMetSerGlyLeuPheLysCysLysVala 372
|||||
1037 GTGCCCTCCCATCCAGCACCGAGACTGGATGAGTGGCAAGAGTTCAATGCAAGGTCA 1056
372 sAnAsnLysAspLeuProAlaProlleGluArgThrIleSerLysProLysGlySerVala 392
|||||
1097 ACAACAAAGAGCTGCCAGCGGCCCTCGAGAGAACCATCTCAAAACCCCAAGGGTCAGTAA 1156
392 tGAlaProGlnValTyrValLeuProProGluGluGluMetThrLysLysGlnValT 412
|||||
1157 GAGCTCCACAGGTATATGTCTGTGCTCCACAGAGAGAGATGCTAAGAAACAGGTCA 1216
412 hrLeuThrCysMetValThrAspPheMetProGluAspIleTyrValGluTyrThrAsnA 432
|||||
1217 CTCTGACCTGCATGGTTCACAGACTTCATGCTGAGACATTTACGTGGAGTGGACCAACA 1276
432 sNGlyLysThrGluLeuAsnTyrLysAsnThrGluProValLeuAspSerAspGlySerT 452
|||||
1277 ACGGGAAACAGAGCTAAACTACAGAAACACTGAACACAGTCTGTGACTCTCATGGTTCTT 1336
452 yrPheMetTyrSerLysLeuArgValGluLysLysAsnTyrValGluArgAsnSerTyrS 472
|||||
1337 ACTTCATGTACAGCAAGCTGAGTGGAAAGAGAGAACTGGTGGAAAGAATAAGTACT 1396
472 erCysSerValHisGluGlyLeuHisAsnHisHisThrThrLysSerPheSerArgT 492
|||||
1397 CTGTGTTAGTGGTCCAGCAGGGTGTGCACAAATCACCACAGCACTAAGAGCTTCTCCCGGA 1456
492 hrProGlyLys 495
|||||
1457 CTCCGGGTAAA 1467

ULT 13

: 9725

AED19725 standard; DNA; 1560 BP.

AED19725;

15-DEC-2005 (first entry)

Anti-PrP antibody heavy chain 44BLH DNA SEQ ID NO 5.

cerebroprotective; cell therapy; gene therapy; drug delivery;

prion disease; 44BLH; ds; therapeutic; monoclonal antibody; heavy chain.

Unidentified.

W02005094846-A1.

13-OCT-2005.

30-MAR-2005; 2005WO-JP006189.

30-MAR-2004; 2004JP-00100649.

(RENO-) RENOMEDIX INST INC.

Fujinaga K, Shinagawa M, Niitsu Y, Hamada H, Horiuchi M;

Homou O, Umetani A;

WPI; 2005-725409/74.

Agent useful for treating prion disease or delivering a substance to a
lesioned region of prion disease, comprises a mesenchymal cell.

Claim 4; SEQ ID NO 5; 34pp; Japanese.

The invention describes an agent (I) for treating prion disease or
delivering a substance to the lesioned region of prion disease,
comprising a mesenchymal cell. Also described are: a nucleic acid (II)
having an anti-prion antibody gene comprising: an antibody heavy chain
gene having SEQ ID No: 1, 3, 5, 30, 32 and 34; a nucleotide sequence
consisting of a degenerate genetic code, which encodes a polypeptide same

CC as that of the above nucleotide sequence, a nucleotide sequence, which is
CC a mutant of the above sequences, or a nucleotide sequence that is
CC complementary to the above sequences and that hybridizes under stringent
CC conditions with the above sequences; and an antibody light chain gene
CC having SEQ ID No: 2, 4, 6, 31, 33 and 35, a nucleotide sequence
CC consisting of degenerate genetic code, which encodes a polypeptide same
CC as that of the above nucleotide sequence, a nucleotide sequence that is
CC a mutant of the above sequences, or a nucleotide sequence that is
CC complementary to the above sequences and that hybridizes under stringent
CC conditions with the above sequences; a vector (III) comprising (II); an
CC anti-prion chimeric antibody (IV) comprising variable region of antibody
CC encoded by (II) and constant region of antibody of animal other than
CC mouse; a nucleic acid that encodes (IV); preparing (MI) a cell having
CC abnormal prion proliferation inhibition activity, comprising transducing
CC a gene that provides abnormal prion proliferation inhibition activity to
CC the cell; a cell (V) having abnormal prion proliferation inhibition
CC activity, being obtainable by (MI) or by utilizing (II) or (III); a
CC sustainable formulation (VI) for discharge of an anti-prion antibody
CC utilized for treating prion disease; use of a mesenchymal cell for
CC producing an agent for delivering a substance to the lesioned region of
CC prion disease; and delivering a substance to the lesioned region of a
CC prion disease, comprising utilizing mesenchymal cell. (I) Is useful for
CC treating prion disease or delivering a substance to the lesioned region
CC of prion disease. (II), (III) Or (MI) is useful for preparing a cell
CC having abnormal prion proliferation inhibition activity. (I), (II),
CC (III), (IV) Or (VI) is useful for treating prion disease. (I) Enables
CC improvement of the symptoms of prion disease. This sequence does not appear in
CC monoclonal antibody heavy chain. Note: This sequence does not appear in
CC the printed specification but has been obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 1560 BP; 413 A; 424 C; 388 G; 335 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	7.55e-104	Length:	1560
Score:	1321.50	Matches:	269
Percent Similarity:	73.7%	Conservative:	19
Best Local Similarity:	68.8%	Mismatches:	39
Query Match:	49.7%	Indels:	65
DB:	14	Gaps:	8

US-10-048-116B-2 (1-495) x AED19725 (1-1560)

Qy 109 AlaThrAsnGluAlaProGlnAlaThrValPheProLysSerProVal-----Leu 125
|||
Dy 560 GCCAAACAAACAGCCCA-----TCGGTCTATCCACTGGCCCTGTGTGGAGGTACA 613
Qy 126 LeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePheProValIleAsn 145
|||
Dy 614 ACTGGCTCCTCGGTGACTTAGGATGCCTGGTCAAGGGTTATTTCCCTGAGCCAGTGACC 673
Qy 146 IleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyrGluThrSerPheLeu 165
:|||||
Dy 674 TTGACCTGG-----AACTCTGGATCCCTGTCAGTGGTGTGCACACCTTCCAGCTCTC 727
Qy 166 ValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPheIleProSerAspAsp 185
:|:|:|
Dy 728 CTGCAGCTCTGAC-----CTCTACACCCCTCAGCAG-CTCAGT----- 762
Qy 186 AspIleTyrAspCysLysValGluHis-TrpGlyLeuGluGluProValLeuLysHisTr 205
:|:|:|
Dy 763 -----GACTGTAACTCGAGCACCTGG----- 784
Qy 205 pGluProGluIleProAlaProMetSerGluLeuThrGluThrGlyGlyGlySerTh 225
Dy 785 -----CCAGCCAGTCCATCACC-----TGCAATGTGGCCCA 816
Qy 225 rThrAlaProSerAlaGlnLeuLysGluLeuGlnAlaLeuGluLysGluAsnAlaGl 245
|||
Dy 817 CCGGCAAGCAGCAGCAACCAAGGTGGCAAGAAAT- 850
Qy 245 nLeuGluTrpGluLeuGlnAlaLeuLysGluLeuAlaGlnAlaAlaSerGluProAr 265
|||||

```

851 -----GAGCCAG 858
265 gGlyProThrIleLysProCysProCysLysCysProAlaProAsnLeuLeuGlyGI 285
859 AGGGCCCAACATCAAGCCCTGCTCATGCAAAATGCCAGCACCTAACTCTTGCGTGG 918
285 yProSerValPheIlePheProProlystIleLysAspValLeuMetIleSerLeuSerPr 305
919 ACATCCGCTTCATCTTCCCTCCAAAGATCAAGGATGACTCATGATCTCCCTGAGCCC 978
305 oileValThrCysValValAspValSerGluAspAspProAspValGlnIleSerTr 325
979 CATAGTCACATGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 1038
325 pPheValAsnValGluValHisThrAlaGlnThrGlnThrHisArgGluAspTyrAs 345
1039 GTTTGTGAACAACGTTGGAAGTACACACAGCTCAGACACAAACCCATAGAGAGATTACAA 1098
345 nSerThrLeuArgValValSerAlaLeuProIleGlnHisGlnAspTyrMetSerGlyLy 365
1099 CAGTACTCTCCGGGTGGTCAAGTGCCTCCCATCCAGCACCCAGGACTGGATGAGTGGCAA 1158
365 sGluPheLysCysLysValAsnAsnLysAspLeuProAlaProIleGluArgThrIleSe 385
1159 GGAGTTCNAATGCAGGTCAACACAAAGACCTCCAGCGCCCATCGAGAGAACCATCTC 1218
385 rLysProLysGlySerValArgAlaProGlnValTyrValLeuProProGluGluGI 405
1219 AAAACCCAAAGGGTCAGTAAGAGTCCACAGGTATATGTCTTGGCTCCACACAGAGNAGA 1278
405 uMetThrLysLysGlnValThrLeuThrCysMetValThrAspPheMetProGluAspTr 425
1279 GATGACTAAGAACAGGTCACTCTGACCTGCATGGTGCACAGACTTCATGCCCTGAAGACAT 1338
425 eTyrValGluTTPThrAsnAsnGlyLysThrGluLeuAsnTyrLysAsnThrGluProVa 445
1339 TTACGTGGAGTGGACCAACACGGGAACACAGAGCTAACTACAGAACACTGACCAGT 1398
445 lIleuAspSerAspGlySerTyrPheMetTyrSerLysLeuArgValGluLysLysAsnTr 465
1399 CCTGCACTCTGATGGTCTTACTTCATGTACAGCAAGCTGAGAGTGGAAAGAACATG 1458
465 pValGluArgAsnSerTyrSerCysSerValValHisGluGlyLeuHisAsnHisHTh 485
1459 GGTGGAAGAAATAGCTACTCTCTGTTCACTGTCCACAGGGGTCTGCACATCACACAC 1518
485 rThrLysSerPheSerArgThrProGlyLys 495
1519 GACTAAGAGCTTCTCCGAGCTCCGGGTAAA 1549

SULT 14
B21727
AEB21727 standard; DNA; 1407 BP.
AEB21727;
08-SEP-2005 (first entry)
Anti-Nogo-antibody 2A10 heavy chain polynucleotide.
neuroprotective; nototropic; cerebroprotective; vasotropic;
antiparkinsonian; anticonvulsant; protein production; therapeutic;
pharmaceutical; amyloidosis; metabolic disorder;
cerebrovascular ischemia; cardiovascular disease; neurological disease;
brain injury; injury; spinal cord injury; vulnerability; dementia;
peripheral neuropathy; parkinson's disease; huntington's chorea;
genetic disorder; Creutzfeldt Jakob disease; infection;
motor neurone disease; cns-gen.; muscular-gen.; myositis;
antiflammatory; inflammation; musculoskeletal disease;
Alzheimers disease; degeneration; antibody; heavy chain; ds.
Mus sp.

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PN WO2005061545-A2.
XX
PD 07-JUL-2005.
XX
PF 20-DEC-2004; 2004WO-GB005343.
XX
PR 22-DEC-2003; 2003GB-00029684.
PR 22-DEC-2003; 2003GB-00029711.
XX
PA (GLAX ) GLAXO GROUP LTD.
XX
PI Hussain I, Prinjha RK,
XX
DR WPI; 2005-522181/53.
XX
FT Modulating production of amyloidogenic peptide in, e.g. Alzheimer's
PT disease, by contacting cell and Nogo polypeptide with Nogo antagonist.
XX
PS Example 4; SEQ ID NO 49; 53pp; English.
XX
CC The invention describes a method of modulating production of an
CC amyloidogenic peptide comprising contacting a cell which is expressing
CC the precursor from which the amyloidogenic peptide is derived and a Nogo
CC polypeptide, with a Nogo antagonist. Also described are: use of a Nogo
CC antagonist in the manufacture of a medicament for the treatment or
CC prophylaxis of a disease involving amyloidosis; and a method of treatment
CC or prophylaxis of Alzheimer's disease comprising administering to the
CC human in need an anti-Nogo antibody. The invention is used for modulating
CC production of amyloidogenic peptide in, e.g. Alzheimer's disease, stroke,
CC traumatic brain injury and spinal cord injury, fronto-temporal dementias,
CC peripheral neuropathy, Parkinson's disease, Huntington's disease,
CC Creutzfeldt-Jakob disease, amyotrophic lateral sclerosis, multiple
CC sclerosis, or inclusion body myositis. The invention provides an
CC unexpected route for therapeutic intervention in particularly Alzheimer's
CC disease. This sequence represents an anti-Nogo-antibody 2A10 heavy chain
CC polynucleotide.
XX
SQ Sequence 1407 BP; 378 A; 396 C; 346 G; 287 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 8.87e-104 Length: 1407
Score: 1320.00 Matches: 285
Percent Similarity: 64.0% Conservative: 26
Best Local Similarity: 58.6% Mismatches: 68
Query Match: 49.6% Indels: 108
DB: 14 Gaps: 13

US-10-048-116B-2 (1-495) x AEB21727 (1-1407)
QY 39 ValTyrGlnSerProGlyAsp-----IleGly----- 47
Db 178 GTGAAGCAGAGGCGCTGGACAAAGCCCTTGATGGATTGGAATATTAACTCTAGCAATGTT 237
QY 48 -----GlnTyrThrHisGluPheAspGlyAspGluLeuPheTyrValAspLeuAspLys 65
Db 238 GGTACTAATAATGAGAAGTTCAAGAGCAAGGCCACACTGACTGTAGACAAATCCTCC 297
QY 66 LysLysThrValTrpArgLeuPro----- 73
Db 298 AGCAGACCTACATGCAGCTCAGCAGCCTGACATCTGAGGACTCTGCGGTCTATTATTGT 357
QY 74 GluPheGlyGlnLeuIleLeuPheGluProGlnGlyGlyLeuGlnAsnIleAlaGlu 93
Db 358 GAACTGGGACAG-----GGCTACTGGGGCCCAAGGCACACTAGTCCCTCAGCC 411
QY 94 LysHisAsnLeuGlyIleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAla 113
Db 412 AAA----- 423
QY 114 ProGlnAlaThrValPheProLysSerProVal-----LeuLeuGlyGlnProAsn 130
Db 424 CCA-----TCGGTCTATCCACTGGCCCTGTGTGTGGAGATACAACTGGCTCCTCGGTG 477

```

131 ThrLeuIleCysPheValAspAsnIlePheProValIleAenIleThrTrpLeuArg 150
|||||
478 ACTCTAGGATCCCTGGTCAAGGGTATTTCCTCAGCCAGTGCCTTGCACCTGG----- 531

151 AsnSerIysSerValThrAspGlyValTyrGluThrSerPheLeuValAsnArgAspHis 170
|||||
532 AACTCTGGATCCCTGCCAGTGGTGCACACACCTTCCAGCTGCCCTGCAGCTCTGCAC--- 588

171 SerPheHisLysLeuSerTyrLeuThrPheIleProSerAspAspAspIleTyrAspCys 190
|||||
589 ---CTTACACCTCAGCAG-CTCAGT-----GACTGT 617

191 LysValGluHis-TrpGlyLeuGluGluProValLeuLysHisTrpGluProGluIlePr 210
|||||
618 AACCTCGAGCACCTGG----- 633

210 oAlaProMetSerGluLeuThrGluThrGlyGlySerThrThrAlaProSerAl 230
|||||
634 ---CCAGCCAGTCCATCACC-----TGCAATGTGGCCACCACCGGCAAGCAGCAC 680

230 aGlnLeuGluLysGluLeuGlnAlaLeuGluLysGluAsnAlaGlnLeuGluTrpGluLe 250
|||||
681 CAAGTGGCAAGAAANT----- 699

250 uGlnAlaLeuGluLysGluLeuAlaGlnAlaSerGluProArgGlyProThrIleLy 270
|||||
700 -----GAGCCACAGGGGCCCAATCAA 722

270 pProCysProProCysLysCysProAlaProAsnLeuLeuGlyGlyProSerValPheIl 290
|||||
723 GCCTGTCTCCATGCAAAATGCCAGCACCTTAACCTCTCTGGGTGGCCCATCCGCTTCAT 782

290 ePheProProLysIleLysAspValLeuMetIleSerLeuSerProIleValThrCysVa 310
|||||
783 CTTCCTCCAAAGATCAAGATGATCTCATGATCTCCTGAGCCCATGATCATCATGTGT 842

310 lValValAspValSerGluAspAspProAspValGlnIleSerTrpPheValAsnAsnVa 330
|||||
843 GGTGGTGGATGTAGCGAGGATGATCCAGATGTCCAGATCAGCTGGTTGTGAAACAACGT 902

330 lGluValHisThrAlaGlnThrGlnThrHisArgGluAspTyrAsnSerThrLeuArgVa 350
|||||
903 GGAAGTACACACAGCTCAGACACAAACCATAGAGAGATTACAACTACTCTCCGGGT 962

350 lValSerAlaLeuProIleGlnHisGlnAspTrpMetSerGlyLysGluPheLysCysLy 370
|||||
963 GGTAGTGCCTCCCTCCATCCAGCACCGAGACTGGATGATGGCAAGAGTTCAATGCNA 1022

370 sValAsnAsnLysAspLeuProAlaProIleGluArgThrIleSerLysProLysGlyse 390
|||||
1023 GGTCAACAACAAGACCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTC 1082

390 rValArgAlaProGlnValTyrValLeuProProGluGluGluMetThrLysLysGl 410
|||||
1083 AGTAAGAGCTCCACAGGTATATGTCTTGCTCCACAGAAGAAGATGACTAAGAAACA 1142

410 nValThrLeuThrCysMetValThrAspPheMetProGluAspIleTyrValGluTrpTh 430
|||||
1143 GGTCACTCTGACCTCGATGGTCAAGACTTTCATGCTGAGACATTTACGTGGAGTGGAC 1202

430 tAsnAsnGlyLysThrGluLeuAsnTyrLysAsnThrGluProValLeuAspSerAspGl 450
|||||
1203 CAACAACGGGAAAACAGAGCTAAACTACAAGAACACTCAACAGCTCTGACTCTGATGG 1262

450 ySerTyrPheMetTyrSerLysLeuArgValGluLysLysAsnTrpValGluArgAsnSe 470
|||||
1263 TTCTTACTTTCATGTACAGCAAGCTGAGAGTGGAAAAGAGAACTGGGTGGAAAGAAATAG 1322

470 rTyrSerCysSerValValHisGluGlyLeuHisAsnHisThrThrLysSerPheSe 490
|||||
1323 CTACTCTCTGTTGATGGTTCACAGGGTCTGCACAACTCAACACAGCACTAAGAGCTTCTC 1382

490 tArgThrProGlyLys 495

Db 1383 CCGACTCCGGTAAA 1398
RESULT 15
AEB08761
ID AEB08761 standard; DNA; 1407 BP.
XX AEB08761;
AC AEB08761;
XX 08-SEP-2005 (first entry)
DT antibody 2A10 heavy chain polynucleotide SEQ ID NO 49.
DE
XX cerebrotective; vasotropic; neuroprotective; vulnerary; nootropic;
XX antiparkinsonian; anticonvulsant; neuroleptic; antibody engineering;
XX pharmacological; cerebrovascular ischemia; cardiovascular disease;
XX neurological disease; brain injury; injury; spinal cord injury;
XX Alzheimers disease; degeneration; dementia; neuropathy;
XX parkinsons disease; Huntingtons chorea; genetic disorder;
XX multiple sclerosis; immune disorder; Creutzfeldt Jakob disease;
XX infection; schizophrenia; psychiatric disorder; motor neurone disease;
XX cns-gen.; muscular-gen.; ds.
OS Unidentified.
XX WO2005061544-A2.
XX 07-JUL-2005.
PD
XX 20-DEC-2004; 2004WO-GB005325.
XX 22-DEC-2003; 2003GB-00029684.
XX 22-DEC-2003; 2003GB-00029711.
PR (GLAX) GLAXO GROUP LTD.
XX
XX Ellis JH, Bon-Duval A, Grundy RI, Hussain F, Mcadam R;
PI Plumpton C, Prinjha RK, Wilson PA;
XX WPI; 2005-479448/48.
DR
XX New antibody or its functional fragment that binds with and neutralizes
XX human neurite outgrowth useful for treating or prophylaxis of stroke and
XX other neurological disease e.g. traumatic brain injury, spinal cord
XX injury, Alzheimer's disease.
XX
XX Example 3; SEQ ID NO 49; 143pp; English.
PS
XX The invention describes an antibody (A1) or its functional fragment, that
XX binds with and neutralizes human neurite outgrowth (NGO). Also described
XX are: providing a first vector encoding a heavy chain of the antibody;
XX providing a second vector encoding a light chain of the antibody; co-
XX transfecting a mammalian host cell with the first and second vectors;
XX culturing the host cell in culture media (preferably serum free) under
XX conditions permissive to the secretion of the antibody from the host cell
XX into the culture media; recovering (and optionally purifying) the
XX secreted antibody; and promoting axonal sprouting involving contacting a
XX human axon with an anti-NGO antibody. The antibody is useful in the
XX preparation of a medicament for treating or prophylaxis of stroke and
XX other neurological diseases/disorders (e.g. traumatic brain injury, spinal
XX cord injury, Alzheimer's disease, frontotemporal dementias (tauopathies),
XX peripheral neuropathy, Parkinson's disease, Huntington's disease and
XX multiple sclerosis); Creutzfeldt-jakob disease (CJD), Schizophrenia,
XX amyotrophic lateral sclerosis (ALS), inclusion body myositis. The
XX antibody inhibits neurodegeneration and/or promotes functional recovery
XX in a human patient suffering, or at risk of developing, stroke or other
XX neurological diseases/disorder. This sequence represents an antibody 2A10
XX heavy chain polynucleotide used in the creation of recombinant anti-NGO
XX antibodies.
XX Sequence 1407 BP; 378 A; 396 C; 346 G; 287 T; 0 U; 0 Other;
SQ Alignment Scores:

ed. No.: 8 87e-104 Length: 1407
ore: 1320.00 Matches: 285
cent Similarity: 64.0% Conservative: 26
st Local Similarity: 58.6% Mismatches: 68
ery Match: 49.6% Indels: 108
14 Gaps: 13

-10-048-116B-2 (1-495) x AEB08761 (1-1407)

39 ValTyrGlnSerProGlyAsp-----ileGly----- 47
178 GTGAAGCAGAGCCCTGGCAAGCCCTTGAGTGGATTGGAAATATTAATCTAGCAATGGT 237
48 -----GlnTyrThrHisGluPheAspGlyAspGluLeuPheTyrValAspLeuAspLys 65
238 GGTACTAATACTAATGAGAAGTTCAAGAGCAAGGCCACACTGACTGTAGACAAATCTCTCC 297
66 LysLysThrValTyrArgLeuPro----- 73
298 AGCAGAGCTACATGACCTCAGCAGCTGACATCTGAGGACTCTGCGGTCTATTATTGT 357
74 GluPheGlyGlnLeuLeuPheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGlu 93
358 GAACCTGGACAG-----GGCTACTCTGGGGCCCAAGGCACACTAGTCACCGTCTCTCTCAGCC 411
94 LysHisAsnLeuGlyIleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAla 113
412 AAA-----ACAAACGCC 423
114 ProGlnAlaThrValPheProLysSerProVal-----LeuLeuGlyGlnProAsn 130
424 CCA-----TCGGTCTATCCACTGCGCCCTGTGTGGAGATACAACTGGCTCTCTCGGTG 477
131 ThrLeuIleCysPheValAspAsnIlePheProProValIleAsnIleThrTrpLeuArg 150
478 ACTTAGGATGCTGCTCAAGGGTATTTCCTGAGCCAGTACCTGACCTGG----- 531
151 AsnSerLysSerValThrAspGlyValTyrGluThrSerPheLeuValAsnArgAspHis 170
532 AACTTGATCCCTCCAGTGGTGTGCACACCTTCCAGCTGTCTCTGAGTCTGAC--- 588
171 SerPheHisLysLeuSerTyrLeuThrPheIleProSerAspAspIleTyrAspCys 190
589 ---CTCTACACCTCAGCAG-CTCAGT-----GACTGT 617
191 LysValGluHis-TrpGlyLeuGluGluProValLeuLysHisTrpGluProGluIlePr 210
618 AACCTCGACACCTGG----- 633
210 oAlaProMetSerGluLeuThrGluThrGlyGlyGlySerThrThrAlaProSerAl 230
634 ----CCAGCCAGTCCATCAC-----TGCAATGTGGCCCAAGCAGCAGCAGC 680
230 aGlnLeuGluLysGluLeuGlnAlaLeuGlnLysGluAsnAlaGlnLeuGluTrpGluLe 250
681 CAAGGTGCACAGAAAT----- 699
250 uGlnAlaLeuGluLysGluLeuAlaGlnAlaSerGluProArgGlyProThrIleLys 270
700 -----GAGCCAGAGGGCCCAATCAA 722
270 sProCysProProCysLysCysProAlaProAsnLeuLeuGlyGlyProSerValPheIle 290
723 GCCCTGTCTCCATGCAATGCCAGCACCTTAACCTCTCTGGGTGGCCCATCGTCTTCAT 782
290 ePheProProLysIleLysAspValLeuMetIleSerLeuSerProIleValThrCysVa 310
783 CTTCCCTCCAAAGATCAAGGATGTACTCATGTATCTCTCTGAGCCCAATAGTCACATGTGT 842
310 lValValAspValSerGluAspAspProAspValGlnIleSerTrpPheValAsnAsnVa 330
843 GGTGTGGATGTGAGCGGAGGATGACCCAGATGTCTCCAGATCAGCTGGTTTGTGAACACAGT 902

QY 330 lgluValHisThrAlaGlnThrGlnThrHisArgGluAspTyrAsnSerThrLeuArgVa 350
Db 903 GGAAGTACACAGCTCAGACACAAACCCATAGAGAGGATTACACAGTACTCTCCGGT 962
QY 350 lValSerAlaLeuProIleGlnHisGlnAspTrpMetSerGlyLysGluPheLysCysLys 370
Db 963 GGTCAAGTCCCTCCCATCCAGCAGCAGGACTGGATGAGTGGCAAGGAGTTCAATGCNA 1022
QY 370 sValAsnAsnLysAspLeuProAlaProIleGluArgThrIleSerLysProLysGlySe 390
Db 1023 GGTCAACAAACAAGACCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTCT 1082
QY 390 rValArgAlaProGlnValTyrValLeuProProGluGluGluMetThrLysLysGln 410
Db 1083 AGTAAGAGCTCCACAGGTATATGTCTTGCTCCACCAGAGAGAGATGACTAAGAAACA 1142
QY 410 nValThrLeuThrCysMetValThrAspPheMetProGluAspIleTyrValGluTrpTh 430
Db 1143 GGTCACTCTGACCTGCATGGTCAAGACTTCATGCTGAAGACATTTACGTGGAGTGGAC 1202
QY 430 rAsnAsnGlyLysThrGluLeuAsnTyrLysAsnThrGluProValLeuAspSerAspGl 450
Db 1203 CAACAAACGGGAAAACAGAGCTAAACTACAAGAAACACTGAACCCAGTCTCTGGACTCTGATGG 1262
QY 450 ySerTyrPheMetTyrSerLysLeuArgValGluLysLysAsnTrpValGluArgAsnSe 470
Db 1263 TTCTTACTTTCATGTACAGCAAGCTGAGAGTGGAAAAGAAAGAACTGGTGGAAAAGAAATAG 1322
QY 470 rTyrSerCysSerValValHisGluGlyLeuHisAsnHisIleThrThrLysSerPheSe 490
Db 1323 CTACTCTCTGTTTCAGTGGTCCAGAGGGTCTGCACAATCACCACAGCACTAAGAGCTTCTC 1382
QY 490 rArgThrProGlyLys 495
Db 1383 CCGGACTCCGGGTAAA 1398

Search completed: May 31, 2006, 23:18:22
Job time : 1013.56 secs

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GenCore version 5.1.8
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protein - nucleic search, using frame_plus_p2n model

on: May 31, 2006, 22:51:28 ; Search time 557.439 Seconds

(without alignments)

5215.692 Million cell updates/sec

le: US-10-048-116B-2_COPY_1_278

fect score: 1496

quence: 1 MPCRSLILGLVLAINTLSI.....QAASBRGPTIKCPCKCF 278

ring table:

BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

ched: 5244920 seqs, 3486124231 residues

al number of hits satisfying chosen parameters: 1049840

imum DB seq length: 0

imum DB seq length: 2000000000

st-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

nmand line parameters:

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=/abs/ABSSWEB spool/US10048116/runat 31052006 110043 25584/app query.fasta 1
3=N Geneseq -QMT-fastap -SUFFIX=p2n.rng -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
NITS=bits -START=1 -END=-1 -MAFIX=blousem62 -TRANSHUMAN40.cdi -LIST=45
CALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
JTFMT=pc -NORM=ext -HEADSIZE=500 -MINLEN=0 -MAXLEN=2000000000 -HOST=abs04
SER=US10048116 @CGN 1.1 942 @runat 31052006 110043 25584 -NCPU=6 -ICPU=3
MMAP -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG -DEV TIMEOUT=120
RN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
APOP=10 -YGAPEXT=0.5 -DLOP=6 -DELEXT=7

abase : N Geneseq 8:*

1: Geneseqn1980s:*

2: Geneseqn1990s:*

3: Geneseqn2000s:*

4: Geneseqn2001as:*

5: Geneseqn2001bs:*

6: Geneseqn2002as:*

7: Geneseqn2002bs:*

8: Geneseqn2003as:*

9: Geneseqn2003bs:*

10: Geneseqn2003cs:*

11: Geneseqn2003ds:*

12: Geneseqn2004as:*

13: Geneseqn2004bs:*

14: Geneseqn2005s:*

15: Geneseqn2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

sult No.	Score	Query Match	Length	ID	Description
1	1496	100.0	1484	5 AAF55098	Aaf55098 DNA encod
2	1214.5	81.2	777	12 ADQ31223	Adq31223 Class II
3	1182	79.0	4713	2 AAV12067	AAV12067 Murine IA

4	1105	73.9	1344	2 AAT60705	Aat60705 cDNA enco
5	1083	72.4	1676	4 ABI99041	Abi99041 Murine pc
6	1078	72.1	776	2 AAQ03169	Aaq03169 Sequence
7	1075	71.9	978	14 ADX26089	Adx26089 Novel cel
8	1072	71.7	776	2 AAT06285	Aat06285 I-Ab-alpha
9	1072	71.7	776	2 AAQ56919	Aaq56919 Mouse I-A
10	1065.5	71.2	1508	2 AAT17587	Aat17587 Vector SC
11	1065.5	71.2	1508	2 AAT86988	Aat86988 SCT1 sing
12	1065.5	71.2	1508	2 AAX89089	Aax89089 Single ch
13	1065.5	71.2	1508	8 ACA60743	ACA60743 Mouse MHC
14	1065	71.2	588	2 AAT60698	Aat60698 Alphalalp
15	1052	70.3	776	2 AAQ35054	Aaq35054 IAB alpha
16	1039	69.5	1013	4 ABI99044	Abi99044 Murine pc
17	1036	69.3	1323	2 AAT60700	Aat60700 cDNA enco
18	1007.5	67.3	1382	2 AAT17588	Aat17588 Vector SC
19	1007.5	67.3	1382	2 AAT86989	Aat86989 SCT1 sing
20	1007.5	67.3	1382	8 ACA60744	ACA60744 Mouse MHC
21	1002	67.0	702	2 AAT60704	Aat60704 cDNA enco
22	996	66.6	588	2 AAT60701	Aat60701 Alphalalp
23	993.5	66.4	1385	2 AAT17586	Aat17586 Vector SS
24	993.5	66.4	1385	2 AAT86987	Aat86987 SSC1 sing
25	993.5	66.4	1385	8 ACA60742	ACA60742 Mouse MHC
26	993	65.7	1243	6 ABN84048	Abn84048 Single ch
27	982	65.6	2053	4 ABI99029	Abi99029 IAS MBP 9
28	982	65.6	2059	4 ABI99032	Abi99032 MBP 1-14
29	949.5	63.5	1701	4 ABI99028	Abi99028 IAS MBP 1
30	949.5	63.5	2346	4 ABI99027	Abi99027 IAS MBP 1
31	946	63.2	2343	4 ABI99033	Abi99033 MBP 90-10
32	944.5	63.1	1662	4 ABI99039	Abi99039 Murine pc
33	944.5	63.1	1680	4 ABI99021	Abi99021 I-As MBP.
34	944.5	63.1	1686	4 ABI99031	Abi99031 MBP 1-14
35	944.5	63.1	1698	4 ABI99038	Abi99038 Murine pc
36	941.5	62.9	1707	4 ABI99030	Abi99030 IAS MBP 9
37	821.5	54.9	768	14 AEB23395	Aeb23395 HLA-DQ al
38	818.5	54.7	768	14 AEB23394	Aeb23394 HLA-DQ al
39	818	54.7	765	14 AEB23392	Aeb23392 HLA-DQ al
40	813.5	54.4	768	6 ABQ74306	Abq74306 Human leu
41	813.5	54.4	768	14 ADV43942	Adv43942 Human psy
42	813.5	54.4	768	14 AEB23385	Aeb23385 HLA-DQ al
43	813.5	54.4	768	14 AEB23386	Aeb23386 HLA-DQ al
44	813.5	54.4	768	14 AEB23393	Aeb23393 HLA-DQ al
45	813.5	54.4	956	4 AAI58351	Aai58351 Human pol

ALIGNMENTS

RESULT 1

AAF55098
ID AAF55098 standard; DNA; 1484 BP.

XX AC AAF55098;

XX DT 15-MAY-2001 (first entry)

XX DE DNA encoding a fusion protein comprising an alpha chain of MHC.

XX KW Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
XX KW major histocompatibility complex; Fc region; antigen; T lymphocyte;
XX KW immunostimulant; vaccine; infection; tumour; ss.

XX OS Synthetic.

XX FH Key Location/Qualifiers

XX CDS 1..1482

XX FT /tag= a

XX PN WO200109194-A1.

XX PD 08-FEB-2001.

XX PF 28-JUL-2000; 2000WO-FR002193.

XX PR 29-JUL-1999; 99FR-00009862.

(CNRS) CNRS CENT NAT RECH SCI.

Glaichenhaus N, Malherbe L;

WPI; 2001-182944/18.

P-PSDB; AAB67480.

New soluble recombinant protein, useful e.g. as immunostimulant.
comprises dimeric major histocompatibility complex molecule fused to
immunoglobulin Fc region.

Example 1; Page 31-33; 43pp; French.

The specification describes soluble recombinant proteins that comprise at least a dimer formed from the alpha and beta-chains of MHC (major histocompatibility complex) Class I and II molecules in which at least one chain has, attached to its C-terminus, at least part of the Fc region of an immunoglobulin. The recombinant proteins, when linked to an antigenic peptide, are used to count and/or purify antigen-reactive T lymphocytes and to characterize their phenotype, e.g. in preclinical evaluation of vaccines. They are also used as immunostimulants, particularly for vaccine development (against infections and tumours), to count and determine phenotype of autoreactive T cells in subjects with, or at risk of developing, autoimmune diseases, e.g. for staging or evaluating treatments, and (to purify and/or enrich Ag-reactive T cells from cell cultures or patient samples, for use in subsequent curative or preventative cellular therapy. The present sequence encodes a recombinant protein of the invention, comprising an alpha chain of MHC molecules

Sequence 1484 BP; 414 A; 394 C; 362 G; 314 T; 0 U; 0 Other;

Comment Scores:
cl. NO.: 7.62e-140 Length: 1484
e: 1496.00 Matches: 278
ent Similarity: 100.0% Conservative: 0
Local Similarity: 100.0% Mismatches: 0
y Match: 100.0% Indels: 0
Gaps: 0

0-048-116B-2_COPY_1_278 (1-278) x AAF55098 (1-1484)

1 MetProCysSerArgAlaLeuLeuLeuGlyValLeuAlaLeuAsnThrMetLeuSerLeu 20
1 ATGCCGTGCAGCAGAGCTCTGATTCCTGGGGTCTCTGCCCTGAACACACCATGCTCAGCCTC 60
21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrThrValTyr 40
61 TCGCGAGGTGAAGACGACATTGAGCGCCGACCGTAGGCTTCTATGGTACAACTGTTTAT 120
41 GlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
121 CAGTCTCTCGAGACATTGGCCAGTACACACATGAATTTGATGGTGATGATGTTCTAT 180
61 ValAspLeuAspLysLysLysThrValTyrArgLeuProGluPheGlyGlnLeuLeu 80
181 GTGGACTTGGATAAGAAGAAACTGTCTGAGGCTTCTCTGAGTTTGGCCAAATTGATATCTC 240
81 PheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGlyLeu 100
241 TTTGAGCCCCAAGGTGGACTGCAAAACATAGCTGCAGAAAAACACAACTTGGGAATCTTG 300
101 ThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPhePro 120
301 ACTAAGAGTCAAAATTCACCCAGCTACCAATGAGGCTCTCTCAGCGACTGTGTCCCC 360
121 LysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePhe 140
361 AAGTCCCTGTGCTGGGTGAGCCCAACACCTTATCTGCTTGTGTGACAACTCTTC 420
141 ProProValIleAsnIleThrTyrArgAsnSerLysSerValThrAspGlyValTyr 160
421 CCACCTGTGATCAACATCATCGGCTCAGAAATAGCAAGTCAGTCACAGCGCGCTTAT 480

QY 161 GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe 180
DB 481 GAGACCACTTCTCTCGTCAACCGTGACCATTCCTTCCACAAAGTCTCTTATCTCACCTTC 540
QY 181 IleProSerAspAspAspIleTyrAspCysLysValGluHisTyrPheGlyLeuGluPro 200
DB 541 ATCCCTCTGATGATGACATTTATGACTGCAAGGTGGAGCACTGGGGCTTGGAGGCGG 600
QY 201 ValLeuLysHisTyrProGluProGluIleProAlaProMetSerGluLeuThrGluThrGly 220
DB 601 GTTCTGAACCACTGGGAACCTGAGATTCCAGCCCCCAATGTCAAGCACTGCAAGAACTGGA 660
QY 221 GlyGlyGlySerThrThrAlaProSerAlaGlnLeuGluLysGluLeuGlnAlaLeuGlu 240
DB 661 GGTGGAGGATCCACTACAGCTCCATCAGCTCGAATAAAGAGCTCCAGGCCCTGGAG 720
QY 241 LysGluAsnAlaGlnLeuGluTyrPheGluGlnAlaLeuGluLysGluLeuAlaGlnAla 260
DB 721 AAGGAAATGCACAGCTCGAAATGGGAGTTGCAAGCACTGGAAGCACTGGCTCAGGCA 780
QY 261 AlaSerGluProArgGlyProThrIleLysProCysProCysProCysLysCysPro 278
DB 781 GCACTGAGCCGAGAGGGCCCAATCAATCAGCCCTGTCTCTCATGCAAAATGCCCA 834

RESULT 2

ADQ31223

ID ADQ31223 standard; cDNA; 777 BP.

XX ADQ31223;

XX ADQ31223;

XX 07-OCT-2004 (first entry)

XX Class II MHC-related I-Ab (alpha)-leucine zipper (LZ) fusion cDNA.

XX class II major histocompatibility complex; MHC; CD4+ T-cell detection;
flow cytometry; mucous membrane invasive antigen;
I-Ab (alpha)-leucine zipper fusion; ss; gene.

OS Unidentified.

XX Key Location/Qualifiers

FT CDS 1..777

FT /*tag= a

FT /product= "Class II MHC-related I-Ab (alpha)-leucine

FT zipper (LZ) fusion protein"

XX JP2004196789-A.

XX 15-JUL-2004.

XX 03-DEC-2003; 2003JP-00404367.

XX 03-DEC-2002; 2002JP-00351818.

XX (SENT-) SENTAN KAGAKU GIJUTSU INCUBATION CENT KK.

XX WPI; 2004-546819/53.

XX P-PSDB; ADQ31222.

XX Peptide-Class II major histocompatibility complex (MHC) composite, useful
for detecting antigen specific CD4+ T-cell, comprises antigen peptide
containing epitope of mucous membrane invasive protein, and extracellular
region of MHC.

XX Example 1; SEQ ID NO 8; 30pp; Japanese.

XX The invention relates to a novel class II major histocompatibility
complex (MHC) antigenic peptide composite comprising a peptide containing
the T-cell antigenic determinant of a mucous membrane invasive protein
and the extracellular region of a class II MHC molecule or at least part
of the extracellular region of the class II MHC molecule having an amino
acid sequence comprising one or more deletions, substitutions or

additions. The molecule of the invention may be useful for detecting an antigen-specific CD4+ T-cell by flow cytometry and for presenting a microorganism-derived mucous membrane invasive protein as an antigen. The method of the invention enables efficient detection of antigen-specific activation of CD4+ T-cells in the mucous membrane. The current sequence is that of the class II major histocompatibility complex-related I-Ab(alpha)-leucine zipper (Lz) fusion cDNA of the invention.

Sequence 777 BP; 193 A; 207 C; 195 G; 182 T; 0 U; 0 Other;

Alignment Scores:

Seq. No.: 5,278-112 Length: 777
 Size: 1214.50 Matches: 232
 Percent Similarity: 92.3% Conservative: 7
 Best Local Similarity: 89.6% Mismatches: 19
 Query Match: 81.2% Indels: 1
 : 12 Gaps: 1

-10-048-116B-2_COPY_1_278 (1-278) x A0Q31223 (1-777)

```

1 MetProCysSerArgAlaLeuLeuLeuGlyValLeuAlaLeuAsnThrMetLeuSerLeu 20
1 ATGCCGCGCAGCAGAGCTCTGATTTCTGGGGGTCTCGCCCTGACCAACCATGCTCAGCCTC 60

21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrcGlyThrThrValTyr 40
61 TGTGGAGGTGAAGACACATTGAGCCGACCATGAGGACCATATGGTATAGTGTATAT 120

41 GlnSerProGlyAspIleGlyGlnThrHisGluPheAspGlyAspGluLeuPheTyr 60
121 CAGTCTCTCTGGAGACATTGCCAGTACACATTGAAATTTGATGGTGATGATGTTCTAT 180

61 ValAspLeuAspIlyslsYsThrValTrpArgLeuProGluPheGlyGlnLeuLeuLeu 80
181 GTGGACTTGGATAAAGAAGAGACTGTCTGGATGCTTCTCGAGTTGGCCAAATTTGGCAAGC 240

81 PheGluProGluGlyGlyLeuGlnAenIleAlaAlaGluLysHisAsnLeuGlyIleLeu 100
241 TTTGACCCCAAGGTGGACTGCAAAACATAGCTGTAGTAAACACAACTTGGAGTCTTG 300

101 ThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPhePro 120
301 ACTAGAGGTCAAATTCACCCAGCTACCAATGAGGCTCTCAAGCGAGTGTGTCCCC 360

121 LysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePhe 140
361 AAGTCCCTGTGCTGTGGGTGAGCCCAACACCTCATCTGCTTTGTGGACAAACATCTTC 420

141 ProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyr 160
421 CTCTCTGTGATCAACATCATATGGCTCAGAAATAGCAAGTCACTGCGACAGCGGTGTAT 480

161 GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe 180
481 GAGACCACTCTTCTCGTCAACGTGACTATCTCTCCCAAGCTGCTTATCTCACTTC 540

181 IleProSerAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGluGluPro 200
541 ATCCCTTTCGACGATGACATTTATGACTCGAAGGTGGAAACACTGGGGCCCTGGAGGAGCG 600

201 ValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGluThrGly 220
601 GTTCTGAAACACTGGGAACCTGAGATTCCAGCCCCCATGTCGCA---CGACACCTGGTT 657

221 GlyGlyGlySerThrThrAlaProSerAlaGlnLeuGluLysGluLeuGlnAlaLeuGlu 240
658 CCGCGCGGATCCACTACAGCTCCATCAGCTCAGCTCGAAGAAAGAGCTCCAGGCCCTGCAG 717

241 LysGluAsnAlaGlnLeuGluTrpGluLeuGlnAlaLeuGluLysGluLeuAlaGln 259
718 AAGGAAATGCAAGCTGGAATGGGAGTTGCAAGCACTGGAAAGGAAGCACTGCTCAG 774

```

AAV12067

ID AAV12067 standard; cDNA; 4713 BP.

XX AAV12067;

XX 08-JUN-1998 (first entry)

XX Murine IAD alpha chain cDNA.

XX Major histocompatibility class II antigen; MHC class II; T cell;

KW T lymphocyte; Th1; Th2; activation; CD4+; antigen presenting cell; APC;

KW autoimmune disease; diabetes; multiple sclerosis; autoimmune thyroiditis;

KW systemic lupus erythematosus; myasthenia gravis; Crohn's disease;

KW inflammatory bowel disease; allergy; asthma; contact sensitivity;

XX immunotherapy; therapy; IAD alpha chain; mouse; ds; circular; cyclic.

OS Mus musculus.

XX WO9746256-A1.

XX 11-DEC-1997.

XX 22-MAY-1997; 97WO-US008697.

XX 23-MAY-1996; 96US-0018175P.

XX (SCRI) SCRIPPS RES INST.

XX Webb SR, Winqvist O, Karlsson L, Jackson MR, Peterson PA;

XX WPI; 1998-041895/04.

XX Synthetic antigen presenting cell for activating CD4+ T cells - useful to treat autoimmune disease, e.g. diabetes, multiple sclerosis, Crohn's disease and inflammatory bowel disease, or allergy, e.g. asthma and contact sensitivity.

XX Example 2; Page 92-94; 141pp; English.

XX This nucleotide sequence comprises a PCR product obtained by amplification of mouse splenocyte cDNA using primers (see AAV12063 and AAV12064) designed for the amplification of IAD alpha chain full-length cDNA. IAD beta chain cDNA (see AAV12068) has been similarly obtained. The IAD sequences were cloned into metallothionein promoter (see AAV12062) - driven vector pRmHs-3 prior to sequencing. Major histocompatibility complex (MHC) class II IAD heterodimers were expressed at the cell surface of transfected Drosophila Schneider 2 (ATCC CRL 10974) cells. The invention relates to the preparation and use of synthetic antigen presenting matrices, in particular antigen presenting cells such as insect cells that have been transfected to produce MHC antigen presenting molecules with one or more accessory molecules. The matrices are used to activate naive CD4+ T cells and to shift the ongoing activation state into a preferred differentiated population of Th1 or Th2 cells. Applications include the treatment of autoimmune disease, e.g. diabetes, multiple sclerosis, autoimmune thyroiditis, systemic lupus erythematosus, myasthenia gravis, Crohn's disease and inflammatory bowel disease, or an allergy, e.g. asthma and contact sensitivity

XX Sequence 4713 BP; 1211 A; 1166 C; 1152 G; 1184 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 1,150-107 Length: 4713
 Score: 1182.00 Matches: 219
 Percent Similarity: 100.0% Conservative: 0
 Best Local Similarity: 100.0% Mismatches: 0
 Query Match: 79.0% Indels: 0
 DB: 2 Gaps: 0

US-10-048-116B-2_COPY_1_278 (1-278) x AAV12067 (1-4713)

QY 1 MetProCysSerArgAlaLeuLeuLeuGlyValLeuAlaLeuAsnThrMetLeuSerLeu 20
 DB 459 ATGCCGCGCAGCAGAGCTCTGATTTCTGGGGGTCTCGCCCTGAAACACCATGCTCAGCCTC 518

21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrValTyr 40
 519 TCGGAGGTGAAGACGACATTTAGGCGCCACACCGTAGGCTTCTATGGTACAACTGTTTAT 578
 41 GlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
 579 CAGTCTCTGGAGACATTTGCCAGGTACACACATGAATTTGATGGTGATGATGTTGTTCTAT 638
 61 ValAspLeuAspLysGlyGlyThrValTrpArgLeuProGluPheGlyGlnLeuIleLeu 80
 639 GTGGACTTGGATAAGAAGAAACTGCTCGAGGCTTCTCGAGGCTTCTGGCCAAATTGATCTC 698
 81 PheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGlyIleLeu 100
 699 TTTGAGCCCCAAGGTGGACTGCAAAAACATAGCTGCAGAAAACACAACTTGGGAACTCTTG 758
 101 ThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPhePro 120
 759 ACTAAGAGGTCAAATTTCAACCCAGCTACCAATGAGGCTCTCTCAAGCGACTGTGTTCCCC 818
 121 LysSerProValLeuLeuGlyGlnProAsnThrIleLeuCysPheValAspAsnIlePhe 140
 819 AAGTCCCTGTGCTGCTGGTGCAGCCCAACACCTTATCTGCTTTGTGGCAACATCTTC 878
 141 ProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyr 160
 879 CCACCTGTGATCAACATCATATGGCTCAGGATAGCAAGTCACTGACAGACGGCGTTTAT 938
 161 GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe 180
 939 GAGACCACTTCTCTGTCGTCACCGTCACCATCTCTTCCCAAGCTGTCTTATCTCACCTTC 998
 181 IleProSerAspAspIleTyrAspCysIysValGluHisTrpGlyLeuGluPro 200
 999 ATCCCTCTGATGATGACATTTATGACTGCAAGGTGGAGCACTGGGGCTGGAGGAGCG 1058
 201 ValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGluThr 219
 1059 GTTCTGAACACTGGGAACCTGAGATTTCAGGCCCCCATGTCTAGAGCTGACAGAACT 1115

'LT 4
' 0705

AAT60705 standard; cDNA; 1344 BP.

AAT60705;

27-AUG-2003 (revised)

12-SEP-1997 (first entry)

cDNA encoding soluble fused MHC heterodimer:peptide complex pLJ23.
 Soluble; fusion; major histocompatibility complex; MHC; heterodimer;
 complex; antigen; binding groove; tolerance; autoantigen; disease;
 insulin dependent; diabetes mellitus; IDDM; antagonist; T cell; anergy;
 presenting cell; de.

Mus sp.

Synthetic.

Key	Location/Qualifiers
mat_peptide	1..1344
	/tag= a

WO9640944-A2.

19-DEC-1996.

07-JUN-1996; 96WO-US010102.

07-JUN-1995; 95US-00480002.

07-JUN-1995; 95US-00482133.

07-JUN-1995; 95US-00483241.

PR 27-OCT-1995; 95US-0005964P.
 XX (ZYMO) ZYMOGENETICS INC.
 PA (ANER-) ANERGEN INC.
 XX Kindsvogel W, Reich EP, Gross JA, Deshpande S, Sheppard PO;
 PI WPI; 1997-052337/05.
 XX P-PSDB; AAW10513.
 DR Novel fused major histocompatibility complex:antigenic peptide complex -
 XX useful to induce tolerance to an autoantigen-related disease e.g. insulin
 PT -dependent diabetes mellitus.
 PT

Example 3; Page 129-132; 142pp; English.

CC The present sequence encodes a novel soluble fused major
 CC histocompatibility complex (MHC) heterodimer:peptide complex, comprising
 CC 1st and 2nd MHC domains, linked by a 5-25 residue linker, an antigenic
 CC peptide able to associate with a peptide binding groove of the MHC
 CC molecule, linked in frame to the 2nd domain by a 5-25 residue linker and
 CC a DNA encoding a 3rd MHC domain linked in frame to the DNA encoding the
 CC antigenic peptide by a DNA encoding a 5-25 residue linker. The complex
 CC can be used to induce immunological tolerance in adults susceptible to,
 CC or suffering from an autoantigen related disease, e.g. insulin dependent
 CC diabetes mellitus (IDDM), by antagonising the binding of particular T
 CC cells and antigen presenting cells, to induce anergy (immunological non-
 CC responsiveness) in the targeted T cell. As the heterodimers and
 CC corresponding antigen are permanently linked into a single chain,
 CC obviating the requirement for complex heterodimer truncation or
 CC formation, the complex eliminates inefficient and non-specific peptide
 CC loading. (Updated on 27-AUG-2003 to correct OS field.)

XX SQ Sequence 1344 BP; 326 A; 356 C; 390 G; 272 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	1.07e-100	Length:	1344
Score:	1105.00	Matches:	203
Percent Similarity:	100.0%	Conservative:	0
Best Local Similarity:	100.0%	Mismatches:	0
Query Match:	73.9%	Indels:	0
DB:	2	Gaps:	0

US-10-048-116B-2_COPY_1_278 (1-278) x AAT60705 (1-1344)

Qy	22	GlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrValTyrGln	41
Db	388	GGTGGCGAGACGACATTTAGGCGCCACCGTAGGCTTCTATGGTACAACTGTTTATCAG	447
Qy	42	SerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyrVal	61
Db	448	TCTCCTGGAGACATTTGGCCAGTACACACATGAATTTGATGGTGTAGTGTGTTCTATGTG	507
Qy	62	AspLeuAspLysGlyThrValTrpArgLeuProGluPheGlyGlnLeuIleLeuPhe	81
Db	508	GACTTGGATAAGAAGAAACTGCTCGAGGCTTCTCGAGGCTTGGCCAAATTGATCTCTTT	567
Qy	82	GluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGlyIleLeuThr	101
Db	568	GAGCCCCAGGTGGAGCTGCANAAACATAGCTGCAGAAAACACAACTTGGGAATCTTGACT	627
Qy	102	LysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPheProLys	121
Db	628	AAGAGGTCAAATTTCAACCCAGCTACCAATGAGGCTCTCAAGCGACTGTGTGTCCCAAG	687
Qy	122	SerProValLeuLeuGlyGlnProAsnThrIleLeuCysPheValAspAsnIlePhePro	141
Db	688	TCCCTGTGCTGGGTGAGCCCAACACCTTATCTGCTTGTGGACACAACTCTTCCCA	747
Qy	142	ProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyrGlu	161
Db	748	CCTGTGATCAACATCACATGGCTCAGAAATAGCAAGTCAGTCACAGACGGCGTTTATGAG	807

922 AATGTGGCCACCGCGCAAGCAGCACCAAGGTGGACAAAGAAATTGAGCCGACGAGGGGCC 981
268 ThrLeuLysProCysProCysProCysLysCysPro 278
982 ACAATCAAGCCCTGTCTCCATGCAAAATGCCCA 1014

LT 6
3169
AAQ03169 standard; DNA; 776 BP.

AAQ03169;
25-MAR-2003 (revised)
31-OCT-2002 (revised)
23-AUG-1990 (first entry)

Sequence encoding the I-Ab-alpha chain of the Class II major histocompatibility complex (MHC) antigens.

I-A (Class II) histocompatibility protein;
major histocompatibility complex antigen; MHC-II; acetylcholine receptor;
myasthenia gravis; autoantigen; autoimmune disease; epitope.

Unidentified.

W08912459-A.
28-DEC-1989.
23-JUN-1989; 89WO-US002784.
23-JUN-1988; 88US-00210594.
21-JUN-1989; 89US-00367751.

(BIOS-) BIOSPAN CORP.

Sharma SD, Lerch BL, Clark BR;

WPI; 1990-022384/03.

New complexes of histo-compatible glycoprotein - with antigenic peptide(s) and label or toxin, used to target antigen specific T helper cells.

Figure 8; Page ?; 74pp; English.

The patent claims complexes of formulae (I), (II) and (III) which are as follows: (i) X - MCH - peptide; (ii) MHC - peptide - X; (iii) MHC - peptide, where X = toxin or labelling gp.; MHC = effective portion of the major histocompatibility glycoprotein; and the peptide includes an epitope associated with one of the major autoimmune diseases. The MHC is the N-terminal of alpha1, alpha2, beta1 or beta2 regions of MHC-II. The complexes can be used to treat and monitor the autoimmune disease. In one protocol, an oligonucleotide encoding the ACHR peptide 195-212 - an epitope in myasthenia gravis patients - was attached to the DNA encoding the N-terminal of the I-Ab-alpha chain. (Updated on 31-OCT-2002 to add missing OS field.) (Updated on 25-MAR-2003 to correct PR field.) (Updated on 25-MAR-2003 to correct PI field.)

Sequence 776 BP; 177 A; 205 C; 199 G; 195 T; 0 U; 0 Other;

Comment Scores:
Cl. No.: 2,56e-98 Length: 776
e: 1078.00 Matches: 201
cent Similarity: 94.1% Conservative: 5
Local Similarity: 91.8% Mismatches: 13
y Match: 72.1% Indels: 0
Gaps: 2

0-048-116B-2_COPY_1_278 (1-278) x AAQ03169 (1-776)

1 MetProCysSerArgAlaLeuLeuGlyValLeuAlaLeuAsnThrMetLeuSerLeu 20
|||||

Db 6 ATGCCGCGCAGCAGAGCTCTGATTCTGGGGGTCTCGCCCTGACACCATGCTCAGCCTC 65
Qy 21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrValTyr 40
Db 66 TGTGAGGTGAAGACGACATTTAGGCCGACCACTAGGCACCTATGGTATTAAGTATAT 125
Qy 41 GlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
Db 126 CAGTCTCTGGAGACATTGGCCAGTACACATTTGAATTTGATGGTGTAGTGTCTTAT 185
Qy 61 ValAspLeuAspLysLysLysThrValTyrArgLeuProGluPheGlyGlnLeuLeu 80
Db 186 GTGGACTTGGATAAGAGGAGACTGTCTGGATGCTTCTGAGTTTGGCCAATTGGCAAGC 245
Qy 81 PheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGlyIleLeu 100
Db 246 TTTGACCCCCAAGGTGGACTGCAAAACATAGCTGTAGTAAACACAACTTGGAGTCTTG 305
Qy 101 ThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPhePro 120
Db 306 ACTAAGAGGTCAAATTCACCCAGCTACCAATGAGGCTCCTCAAGCGACTGTGTTCCTC 365
Qy 121 LysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePhe 140
Db 366 AAGTCCCTGTGCTGGGTGAGCCCAACCCCTCATCTGCTTTGTGGACAACTCTTC 425
Qy 141 ProProValIleAsnIleThrTyrLeuArgAsnSerLysSerValThrAspGlyValTyr 160
Db 426 CCTCTGTGATCAACATCAGATGCTCAGAAATAGCAAGTCAGTCGACAGCGTCTTAT 485
Qy 161 GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe 180
Db 486 GAGACAGCTTCTTCGTCAACCGTACTATTCCTTCCACAAGCTGCTTATCTACCTTC 545
Qy 181 IleProSerAspAspAspIleTyrAspCysLysValGluHisTyrGlyLeuGluPro 200
Db 546 ATCCCTTCTGACGATGACATTTATGACTGCAAGGTGGAAACACTGGGGCTTGGAGGAGCG 605
Qy 201 ValLeuLysHisTyrGluProGluIleProAlaProMetSerGluLeuThrGluThr 219
Db 606 GTTCTGAACACTGGGAACCTGAGATTCACGCCCCCATGTCAGAGCTGACAGAGACT 662

RESULT 7

ADX26089
ID ADX26089 standard; DNA; 978 BP.

AC ADX26089;

DT 05-MAY-2005 (first entry)

DE Novel cell pain response detection method-related mouse gene SeqID435.

KW pain; animal disease model; expression; analgesic; antiaddictive;
KW nootropic; anticonvulsant; vasotropic; neuroprotective; tranquilizer;
KW antiasthmatic; antirheumatic; antiarthritic; osteopathic;
KW ophthalmologic; antiinflammatory; antipruritic; dermatological;
KW antileuc; gastrointestinal-Gen.; nephrotoxic; gynecological;
KW hepatotropic; antiparkinsonian; neuroleptic; laxative; gene therapy;
KW neuropathic pain; Alzheimers disease; Parkinsons disease;
KW motor neurone disease; Huntingtons disease; schizophrenia; gene; ds.

OS Mus sp.

PN WO2005014849-A2.

PD 17-FEB-2005.

PF 06-JUL-2004; 2004WO-US023166.

PR 03-JUL-2003; 2003US-0485101P.

PA (EURO-) EUROCELTIQUE SA.

XX

Tong J, Jin G, Ji R, Xu Y, Chiang LW, Lavery DU;

WPI; 2005-163258/17.

Detecting pain responses in a cell, useful in identifying potential therapeutic and diagnostic candidates for treating pain, by identifying genes that are differentially expressed in a model of neuropathic pain.

Example 1; SEQ ID NO 435; 173pp; English.

This invention relates to a novel method of detecting a pain response in a cell which comprises determining the expression level in a test cell of at least one nucleic acid molecule and comparing the expression level to a level in an animal model of pain, where similar or identical expression levels indicate a pain response in the test cell. The invention may be useful for the development of compounds with an analgesic, antiaddictive, neurotropic, anticonvulsant, vasotropic, neuroprotective, tranquilizer, antiasthmatic, antirheumatic, antiarthritic, osteopathic, ophthalmological, antiinflammatory, antipruritic, dermatological, antitumor, gastrointestinal, antipruritic, antipruritic, dermatological, hepatotropic, antiparkinsonian, neuroleptic or laxative activity whilst the disclosed sequences may prove useful for gene therapy. The methods and compositions of the present invention are useful for identifying agonists and antagonists for the gene or gene products as potential therapeutic and diagnostic candidates for treating pain, including neuropathic pain, nociceptive pain, chronic pain, inflammatory pain, pain associated with cancer, and pain associated with a rheumatic disease, and also for addiction, seizure, stroke, ischemia, a neurodegenerative disorder, anxiety, depression, headache, asthma, rheumatic disease, osteoarthritis, retinopathy, inflammatory eye disorders, pruritus, ulcer, gastric lesions, uncontrollable urination, an inflammatory or unstable bladder disorder, inflammatory bowel disease, irritable bowel syndrome (IBS), irritable bowel disease (IBD), gastroesophageal reflux disease (GERD), functional dyspepsia, functional chest pain of presumed esophageal origin, functional dysphagia, non-cardiac chest pain, symptomatic gastroesophageal disease, gastritis, aerophagia, functional constipation, functional diarrhea, borborygmi, chronic functional abdominal pain, recurrent abdominal pain (RAP), function abdominal bloating, functional biliary pain, functional incontinence, functional ano-rectal disorder, cholecystitis, interstitial cystitis, dysmenorrhea, or dyspareunia. They can also be used for diagnosing or treating Alzheimer's disease, Parkinson's disease, amyotrophic lateral sclerosis, Huntington's disease and schizophrenia. The present sequence is that of a mouse gene which demonstrated homology to a human gene whose expression level analysed in the method of the invention.

Sequence 978 BP; 218 A; 263 C; 247 G; 250 T; 0 U; 0 Other;

Alignment Scores:

Seq. No.:	7.05e-98	Length:	978
Score:	1075.00	Matches:	203
Percent Similarity:	93.6%	Conservative:	2
Local Similarity:	92.7%	Mismatches:	14
Match:	71.9%	Indels:	0
	14	Gaps:	0

-10-048-116b-2_COPY_1_278 (1-278) x ADX26089 (1-978)

1 MetProCysSerArgAlaLeuLeuLeuValLeuValLeuAlaLeuAsnThrMetLeuSerLeu 20
24 ATGCCGCGCAGACAGCTCTGATTCCTGGGGTCTCGCCCTGACCATGCTCAGCCTC 83
21 CysGlyGlyGluAspAspLeuGluAlaAspHisValGlyPheThrGlyThrThrValTyr 40
84 TGGCGAGGTGAACAGACATTGAGCGCGACACGATAGGCTCTTATGATTAAGTATAT 143
41 GlnSerProGlyAspLeuGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
144 CAGTCCTCGGAGACATTGGCCAGTACACATTGAAATTTGATGATGATGATGTTCTAT 203
61 ValAspLeuAspLeuValThrValTyrArgLeuProGluPheGlyGlnLeuLeu 80
204 GTGGACTTGGATTAAGAGGAGACTGCTGGATGCTTCTCGAGTTTCTCACTGAGAAGA 263

QY	81	PheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisLeuLeuGlyIleLeu	100
DB	264	TTTGAGCCCCAAGGTGGACTGCAAAACATAGCTACAGGAAACACAACTTGGAAATCTTG	323
QY	101	ThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPhePro	120
DB	324	ACTNAGAGGTCAAATTCACCCAGCTACCAATGAGGCTCCTCAGGAGCTGTGTTCCTCC	383
QY	121	LysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePhe	140
DB	384	AAAGTCTCTCTGCTGCTGGGTGAGCCCAACACCTTATCTGCTTGTGGCAACATCTTC	443
QY	141	ProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyr	160
DB	444	CTCTCTGTGATCAACATCATGCTGCTCAGAAATAGCAAGTCACTCAGACGCGGTTAT	503
QY	161	GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe	180
DB	504	GAACACGCTTCTTCTGCTCAACCGTGACTATTCTTCCACAGCTGTCTTATCTCACCTTC	563
QY	181	IleProSerAspAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGluPro	200
DB	564	ATCCCTTCTGACGATGACATTTATGACTGCAAGGTGGAGCACTGGGGCTTGGAGGAGCG	623
QY	201	ValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGluThr	219
DB	624	GTTCGAAACACCTGGGAACCTGAGATTCCAGCCCCCATGTCCAGAGCTGACAGAGACT	680
RESULT	8		
ID	AAT06285		
XX	AAT06285 standard; DNA; 776 BP.		
AC	AAT06285;		
XX	10-APR-1996 (first entry)		
DT	1-AB-alpha chain DNA.		
DE	I-Ab-alpha-chain; acetylcholine receptor; myelin basic protein;		
KW	autoantigen; MHC class II; major histocompatibility complex;		
KW	autoimmunity; autoimmune disease; rheumatoid arthritis;		
KW	myasthenia gravis; multiple sclerosis; allograft rejection; vaccine; ds.		
XX	Mus musculus.		
XX	US5468481-A.		
XX	21-NOV-1995.		
XX	14-APR-1992; 92US-00869293.		
XX	23-JUN-1988; 88US-00210594.		
XX	21-JUN-1989; 89US-00367751.		
XX	30-AUG-1990; 90US-00576084.		
XX	28-DEC-1990; 90US-00635840.		
XX	23-APR-1991; 91US-00690840.		
XX	(ANER-) ANERGEN INC.		
XX	Sharma SD, Lerch BL, Clark BR;		
XX	WPI; 1993-036056/04.		
XX	Pure major MHC-peptide complex - useful in treating deleterious immune		
XX	response such as autoimmunity.		
XX	Disclosure; Fig 8; 47pp; English.		
XX	A sequence (AAT06285) encoding the I-Ab-alpha chain is utilised in the		
XX	construction of novel MHC class II conjugates. An autoantigen peptide,		
XX	derived e.g. from the acetylcholine receptor alpha chain (AAR86421) or		
XX	myelin basic protein (AAR86422), is attached to the N-terminal end of an		

141 ProProValIleAenIleThrTrpLeuAtcAAsnSerIysSerValThrAspGlyValTyr 160
 426 CTTCTGTGTATCAACATCATGCTCTAGAAAGCAAGTCAGTCGACGCGTGTAT 485
 161 GluThrSerPheLeuValAenArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe 180
 486 GAGACCGAGCTTCTTCGTCACCGTGACTATTCTCTCCACAGCTGTCTTATCTCACCTTC 545
 181 IleProSerAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGluGluPro 200
 546 ATCCCTCTGACGATGACATTTATGACTGCAAGGTGGAACACTGGGGCTTGGAGGACCG 605
 201 ValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGluThr 219
 606 GTTCTGAACACTGGGAACCTGAGATTCACGCCCTCATGTTCAGAGCTGACAGAGACT 662

SULT 10

T17587

AAT17587 standard; DNA; 1508 BP.

AAT17587;

26-SBP-1996 (first entry)

Vector SCT1-derived single chain gene encoding MHC fusion complex.

MHC; major histocompatibility complex; PCR; polymerase chain reaction;
 T cell activity modulator; antagonist; immune disorder; allergy;
 multiple sclerosis; insulin-dependent diabetes mellitus;
 rheumatoid arthritis; myasthenia gravis; ds.

Synthetic.

Key Location/Qualifiers
 CDS 6..1508
 /tag= a
 sig_peptide 6..86
 /tag= b
 /label= I-Ad beta chain leader
 /note= "murine MHC class II I-Ad gene beta chain leader sequence"
 misc_feature 87..137
 /tag= c
 /label= OVA_323-339
 /note= "chicken ovalbumin residues 323-339"
 misc_feature 138..167
 /tag= d
 /note= "10 residue linker peptide"
 misc_feature 168..452
 /tag= e
 /label= I-Ad beta1
 /note= "murine MHC class II I-Ad gene beta-1 domain"
 misc_feature 453..734
 /tag= f
 /label= I-Ad beta2
 /note= "murine MHC class II I-Ad gene beta-2 domain"
 misc_feature 735..806
 /tag= g
 /note= "24 residue peptide linker"
 misc_feature 807..1067
 /tag= h
 /label= I-Ad alpha1
 /note= "murine MHC class II I-Ad gene alpha-2 domain"
 misc_feature 1068..1352
 /tag= i
 /label= I-Ad alpha2
 /note= "murine MHC class II I-Ad gene alpha-2 domain"
 misc_feature 1353..1505
 /tag= j
 /label= I-Ad alpha-TM
 /note= "murine MHC class II I-Ad gene alpha-transmembrane domain"

XX WO9604314-A1.
 PD 15-FEB-1996.
 XX 31-JUL-1995; 95WO-US0009816.
 XX 29-JUL-1994; 94US-00283302.
 PR 01-FEB-1995; 95US-00382454.
 XX (DADE-) DADE INT INC.
 XX Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
 PI Chavalliaz P, Jiao J;
 XX WPI; 1996-129343/13.
 DR P-PSDB; AAR98906.
 XX
 PT Major histocompatibility complex fusion complex for modulating T cell
 PT activity - used in the treatment of immune disorders, e.g. multiple
 PT sclerosis, IDDM and rheumatoid arthritis.
 XX
 PS Example 17; Fig 28; 210pp; English.
 XX
 CC AAT17587 encodes a murine MHC fusion complex capable of modulating T cell
 CC activity encoded by the vector SCT1. The MHC fusion complex comprises at
 CC least one MHC molecule containing a peptide-binding groove and a
 CC presenting peptide covalently linked to the MHC molecule and opt. a
 CC transmembrane domain. DNA encoding a MHC fusion complex may be cloned
 CC into a host cell to express the complex. The transformed cells may then
 CC be used to identify peptides that modulate, pref. antagonise, T cell
 CC activity. DNA encoding a MHC fusion complex or a single chain fusion
 CC molecule may be used to vaccinate a mammal against a targeted disorder.
 CC The fusion complexes may be used to suppress an immune response in an
 CC animal suffering from an immune disorder e.g. multiple sclerosis, insulin
 CC -dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or
 CC chronic allergies. The complexes may also be used in the treatment of
 CC livestock and pets such as cats and dogs. The MHC fusion complexes can be
 CC produced such that they contain a single antigenic peptide including one
 CC of known structure, additionally a wide range of peptides can be
 CC presented for T cell interaction
 XX
 SQ Sequence 1508 BP; 337 A; 414 C; 440 G; 317 T; 0 U; 0 Other;
 Alignment Scores:
 Pred. No.: 1.15e-96 Length: 1508
 Score: 1065.50 Matches: 198
 Percent Similarity: 98.5% Conservative: 0
 Best Local Similarity: 98.5% Mismatches: 0
 Query Match: 71.2% Indels: 3
 DB: 2 Gaps: 1
 US-10-048-116B-2_COPY_1_278 (1-278) x AAT17587 (1-1508)
 QY 22 GlyGly-----GluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrThr 38
 Db 792 GCGGGTTCCTCGAGTGAAGACGACATTGAGCGCAGCACAGTAGGCTTCTATGTACTAACT 851
 QY 39 ValTyrGlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeu 58
 Db 852 GTTTATCAGTCTCCTGGAGACATTGGCCAGTACACACATGATTTGATGATGATGTTG 911
 QY 59 PheTyrValAspLeuAspLysLysThrValTrpArgLeuProGluPheGlyGlnLeu 78
 Db 912 TTCTATGTGAGCTTGGATAAGAGAAACTGTCTGGAGGCTTCTCGAGTTTGGCCAAATTG 971
 QY 79 IleLeuPheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisGlnGly 98
 Db 972 ATACTCTTTGAGCCCCCAGGTGAGCTGCAGAAACATAGCTGCAGAAAACACAACTTGGGA 1031
 QY 99 IleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnIleThrVal 118
 Db 1032 ATCTTGACTAAGAGGTCAAAATTTACCCCGAGCTTACCAATGAGGCTCCTCAAGCGACTGTG 1091

119 pheProLysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAen 138
|||||
1092 TTCCCAAGTCCCTGCTGCTGGGTGAGCCCAACACCCCTTATCTGTTGTGGACAA 1151
|||||
139 llePheProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGly 158
1152 ATCTTCCCACTGTGATCAACATCATGGCTCAGAAATAGCAAGTCAAGTCAACACGGC 1211
|||||
159 ValTyrGluThrSerPheLeuValIleAsnArgAspHisSerPheHisLysLeuSerTyrLeu 178
1212 GTTTATGAGACCAAGTCTCTGCTCAACCGTGACCATTCCTCCCAAGCTGCTTATCTC 1271
|||||
179 ThrPheIleProSerAspAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGlu 198
1272 ACCTTCATCCCTTCTGATGATGACATTTATGACTGCNAGGTGGAGCACTGGGGCTGGAG 1331
|||||
199 GluProValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGlu 218
1332 GAGCGGTTCTGAACACACTGGGAACCTGAGATTCCAGCCCCCATGTCTCAGAGCTGACAGAA 1391
219 Thr 219
1392 ACT 1394

LT 11

6988

AAT86988 standard; DNA; 1508 BP.

AAT86988;

27-MAR-1998 (first entry)

SC1 single chain gene.

Construction; major histocompatibility complex; MHC; fusion complex;

SC1 single chain gene; ss.

Synthetic.

Key Location/Qualifiers
CDS 6..1508
/*tag= a

WO9728191-Al.

07-AUG-1997.

30-JAN-1997; 97WO-US001617.

31-JAN-1996; 96US-00596387.

(DADE-) DADE INT INC.

Rhode PR, Jiao J, Burkhardt M, Wong HC;

WPI: 1997-402555/37.

P-PSDB; AAW29213.

Single chain major histocompatibility complex comprising linked alpha and beta chains - useful for suppressing an immune response to an auto-immune disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes mellitus, etc.

Example 17; Page 137-139; 217pp; English.

The present sequence was used in the construction of major histocompatibility complex (MHC) fusion complexes

Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

ment Scores:

ti. No.:

1.15e-96

Length:

1508

Score: 1065.50 Matches: 198
Percent Similarity: 98.5% Conservative: 0
Best Local Similarity: 98.5% Mismatches: 0
Query Match: 71.2% Indels: 3
DB: 2 Gaps: 1

US-10-048-116b-2_COPY_1_278 (1-278) x AAT86988 (1-1508)

QY 22 GlyGly-----GluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrThr 38
|||||
Db 792 GCGGTTCTCGAGTGAAGACGACATTGAGCCCGACAGTAGGCTTCTATGTCACACT 851
|||||
QY 39 ValTyrGlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeu 58
852 GTTTATCAGTCTCCTGGAGACATTGGCCAGTACACATGAATTTGATGTTGATGAGTTG 911
|||||
QY 59 PheTyrValAspLeuAspLysLysLysValTyrValTyrArgLeuProGluPheGlyGlnLeu 78
912 TTCTATGTGGACTTGGATAAGAAGAAACTGTCTGGAGGCTTCTGAGTTTGGCCATTTG 971
|||||
QY 79 IleLeuPheGluProGlnGlyGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGly 98
972 ATACTTTTGAGCCCAAGGTGGACTGCAGAAACATAGCTGCAGAAAACACAACTTGGCA 1031
|||||
QY 99 IleLeuThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrVal 118
1032 ATCTTGACTAAGAGGTCAAAATTTACCCCCAGCTACCAATGAGGCTCTCAAGCGACTGTG 1091
|||||
QY 119 PheProLysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAen 138
1092 TTCCCAAGTCCCTGCTGCTGGGTGAGCCCAACACCCCTTATCTGTTGTGGACAAAC 1151
|||||
QY 139 IlePheProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGly 158
1152 ATCTTCCCACTGTGATCAACATCATGGCTCAGAAATAGCAAGTCAAGTCAACACGGC 1211
|||||
QY 159 ValTyrGluThrSerPheLeuValIleAsnArgAspHisSerPheHisLysLeuSerTyrLeu 178
1212 GTTTATGAGACCAAGTCTCTGCTCAACCGTGACCATTCCTCCCAAGCTGCTTATCTC 1271
|||||
QY 179 ThrPheIleProSerAspAspAspIleTyrAspCysLysValGluHisTrpGlyLeuGlu 198
1272 ACCTTCATCCCTTCTGATGATGACATTTATGACTGCNAGGTGGAGCACTGGGGCTGGAG 1331
|||||
QY 199 GluProValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGlu 218
1332 GAGCGGTTCTGAACACACTGGGAACCTGAGATTCCAGCCCCCATGTCTCAGAGCTGACAGAA 1391
219 Thr 219
1392 ACT 1394

RESULT 12

AA89069

ID AA89069 standard; DNA; 1508 BP.

XX

XX AA89069;

XX 14-SEP-1999 (first entry)

XX Single chain IAD/OVA 323-229 MHC fusion protein encoding DNA.

DE Major histocompatibility complex; MHC; single chain MHC; sc-MHC; Ig;
KW peptide binding groove; immunoglobulin; T cell receptor; immune response;
KW immune-related disorder; antigenic peptide; fusion protein; ss.

XX Synthetic.

OS WO9921572-Al.

PN 06-MAY-1999.

PD 13-OCT-1998; 98WO-US021520.

XX

PF

29-OCT-1997; 97US-00960190.

(SUNO-) SUNOL MOLECULAR CORP.

Rhode PR, Acevedo J, Burkhardt M, Jiao J, Wong HC;

WPI; 1999-418411/35.

P-PSDB; AAY27111.

Single chain major histocompatibility complex class I complexes.

Example 1; Fig 1; 148pp; English.

The invention relates to new single chain major histocompatibility complex (sc-MHC) class II complexes that comprise a peptide binding groove, and a modified class II beta 2 chain or covalently linked immunoglobulin (Ig) light chain constant (C1) region. The MHC complexes are useful for detection and analysis of peptide ligands, pathogenic T-cells, for functional, cellular and molecular assays. They can be used to identify and isolate T cell receptor and/or MHC agonists and antagonists. They can be used in vivo to compete with pathogenic antigen presenting cells involved in immune-related disorders. They can also be used to raise antibodies and to screen immune cells. It is also use in a method of suppressing an immune response in mammals. The sc-MHC complexes comprising modified class II beta 2 chains and/or Ig-C1 regions are soluble and provide enhanced yield. These MHC complexes also can contain single antigenic peptides readily isolated from expressing cells in significant quantities. The polyspecific MHC complexes also provide a means to detect cells expressing multiple target structures with a single complex. The present sequence represents a DNA encoding a single chain IAd/OVA 323-229 MHC fusion protein

Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

Alignm Scores:

Seq. No.:	1.15e-96	Length:	1508
Score:	1085.50	Matches:	198
Percent Similarity:	98.5%	Conservative:	0
Identical Similarity:	98.5%	Mismatches:	0
Indels:	3		
Gaps:	1		

-10-048-116B-2_COPY_1_278 (1-278) x AAX89069 (1-1508)

```
22  GlyGly-----GluAspAspIleGluAlaAspHisValGlyPheTyGlyThrThr 38
792  GCGGGTCTCTCGAGTGAAGACGACATTGAGCGCGACCGTAGGCTTCTATGTCACAACT 851
39  ValTyGlnSerProGlyAspIleGlyGlnTyThrHisGluPheAspGlyAspGluLeu 59
852  GTTTATCAGTCTCTCGAGACATTGGCGCAGTACACACATGAAATTGATGTCATGAGTTG 911
59  PheTyValAspLeuAspIleGlyValTyThrValTyArgLeuProGluPheGlnLeu 78
912  TTCTATGTGACTTGATAGAGAAACTGCTGAGGGCTTCTGAGTTTGCCCAATTG 971
79  IleLeuPheGluProGlnGlyLeuGlnAsnIleAlaAlaGluTyHisAsnLeuGly 98
972  ATACTCTTTGAGCCCGAGGTGACCTGCAGAAACATAGCTGCAGAAACACAACTTGGGA 1031
99  IleLeuThrTyArgSerAsnAspThrProAlaThrAsnGluAlaProGlnAlaThrVal 118
1032  ATCTTGACTAAGAGGTCAAAATTTTCCGCCAGCTACCAATGAGGCTCTCAAGCGACTGTG 1091
119  PheProTySerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsn 138
1092  TTCCCCAAGTCCCTGTGCTGTGGGTGACGCCCAACACCCCTTATCTGCTTGTGGCAAC 1151
139  IlePheProValIleAsnIleThrTyLeuArgAsnSerTySerValThrAspGly 158
1152  ATCTTCCCACTGTGATCAACATCAATCATGCTCAGAAATAGCAAGTCAGTCACAGAGCGC 1211
```

```
QY 159 ValTyGluThrSerPheLeuValAsnArgAspHisSerPheHisLeuSerTyLeu 178
Db 1212 GTTATGAGACCGAGCTTCTCTGTCACCGTGACCATCTTCCACAAGCTGTCTATCTC 1271
QY 179 ThrPheIleProSerAspAspIleTyArgPheCysValGluHisTrpGlyLeuGlu 198
Db 1272 ACCTTCATCCCTTCTGATGATGACATTATGACTGCAAGGTGAGCACTGGGCGCTGGAG 1331
QY 199 GluProValLeuTyHisTrpGluProGluIleProAlaProMetSerGluLeuThrGlu 218
Db 1332 GAGCCGGTCTGAAACACTGGGAACCTGAGATTCCAGCCCCCATGTGACAGCTGACAGAA 1391
QY 219 Thr 219
Db 1392 ACT 1394
RESULT 13
ACA60743
ID ACA60743 standard; DNA; 1508 BP.
XX ACA60743;
XX
DT 16-JUN-2003 (first entry)
DE Mouse MHC I-Ad/Ova 323-339 synthetic gene SCTL.
XX
KW MHC; major histocompatibility complex; gene therapy; fusion complex;
KW peptide-binding groove; T cell modulation; class II MHC; vaccine;
KW autoimmune disorder; multiple sclerosis; rheumatoid arthritis;
KW insulin-dependent diabetes mellitus; myasthenia gravis; immunogen;
KW chronic allergy; mouse; ds; I-Ad; gene.
XX
OS Mus sp.
XX Synthetic.
XX
PN US2002198144-A1.
PD 26-DEC-2002.
PF 06-JUL-2001; 2001US-00900379.
PR 29-JUL-1994; 94US-00283302.
PR 01-FEB-1995; 95US-00382454.
PR 17-JAN-1997; 97US-00776084.
XX
PA (DADE-) DADE INT INC.
XX
PI Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
PI Chavalliaz P, Jiao JJ;
XX
XX WPI; 2003-341126/32.
DR P-PSDB; ABU72107.
XX
PT Novel major histocompatibility complex fusion complex having presenting
PT peptide covalently linked to MHC molecule containing peptide-binding
PT groove, used for suppressing immune response in multiple sclerosis,
PT allergies.
XX
PS Example 17; Fig 28; 126pp; English.
XX
CC The invention relates to a major histocompatibility complex (MHC) fusion
CC complex (I) comprising an MHC molecule that contains a peptide-binding
CC groove, and a presenting peptide covalently (e.g. an antigenic peptide)
CC linked to the MHC molecule, where (I) is capable of modulating the
CC activity of a T cell. Also included are a DNA construct coding for the
CC complex, where the MHC molecule is a class II MHC (e.g. mouse I-Ad or I-
CC As, or human HLA-DR1 (human leukocyte antigen-DR1)), a multivalent MHC
CC fusion complex comprising two or more linked complexes, identifying a
CC peptide that can modulate the activity of T cells (involving introducing
CC into host cells cloning vectors that each contain the fusion complex DNA,
CC culturing the host cells under conditions suitable for expression of the
CC MHC fusion complex, and selecting host cells that express MHC fusion
CC complex that modulate the activity of T cells), a single recombinant
```

expression vector comprising DNA that codes for the alpha and beta chains of the fusion complex MHC protein, a single recombinant expression vector comprising DNA that codes for a T cell costimulatory factor and the alpha and beta chains of the MHC fusion complex. The DNA constructs can contain heterologous leader peptide sequences and Kozak sequence for efficient expression of the fusion complex. Also included are inducing an immune response in a mammal (including vaccinating a mammal against a targeted disorder, by administering DNA sequence comprising a fusion complex, or DNA sequence coding for a fusion complex which is a single chain fusion molecule) and suppressing an immune response in a mammal by administering to the mammal a DNA sequence comprising an expression vector, encoding a full length MHC molecule that contains a transmembrane domain, and a presenting peptide that is a T cell receptor (TCR) antagonist or partial agonist and is covalently linked to the MHC protein, or DNA sequence coding for the fusion complex which is a single chain fusion molecule. The methods are useful for identifying a peptide that can modulate the activity of T cells, inducing an immune response in a mammal (including vaccinating a mammal against a targeted disorder) and for suppressing an immune response in a mammal. The disorders include an autoimmune disorder such as multiple sclerosis, insulin-dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or chronic allergies. The present sequence encodes a mouse MHC class II I-Ad fusion complex of the invention

Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

nment Scores:			
e:	1.15e-96	Length:	1508
ent Similarity:	1065.50	Matches:	198
Local Similarity:	98.5%	Conservative:	0
y Match:	98.5%	Mismatches:	0
	71.2%	Indels:	3
	8	Gaps:	1

0-048-116B-2_COPY_1_278 (1-278) x ACA60743 (1-1508)

22	GlyGly-----	GlulAspIleGluAlaAspHisValGlyPheTyrGlyThrThr	38
792	GGCGGTTCTCGAGTGAAGACGACATTTAGGCGGACCATGAGGCTTCTATGGTACAACT		851
39	ValTyrGlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeu		58
852	GTTTATCAGTCTCTCGAGACATTTGGCCAGTACACACATGAAATTTGATGGTATGAGTTG		911
59	PheTyrValAspLeuAspLysLysThrValTyrPheGluPheGlyGlnLeu		78
912	TTCTATGGACTTGGATGAAGAAACTGCTGGAGGCTTCTGAGTTTGGCCAAATG		971
79	IleLeuPheGluProGlnGlyGlyLeuGlnAenlleAlaAlaGluLysHisAsnLeuGly		98
972	ATATCTTTGAGCCCAAGGTGGACTGCAAAACATAGCTGCAGAAAAACACAACTTGGGA		1031
99	IleLeuThrLysArgSerAsnThrProAlaThrAsnGluAlaProGlnAlaThrVal		118
1032	ATCTTGACTAAGAGGTCAAATTTACCCAGCTACCAATGAGGCTCTCAAGCGACTGTG		1091
119	PheProLysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAen		138
1092	TTCCCAAGTCCCTGTGCTGGTGGTGCAGCCACACACCTTATCTGCTTGTGGACAAC		1151
139	IlePheProProValIleAenlleThrTrpLeuArgAsnSerLysSerValThrAspGly		158
1152	ATCTTCCCACTGTGATCAACATCATGGCTCAGAAATAGCAAGTCAGTCACAGACGGC		1211
159	ValTyrGluThrSerPheLeuValAenArgAspHisSerPheHisLysLeuSerTyrLeu		178
1212	GTTTATGAGACCAAGCTTCTCTGCTCAACCGTGACCATTTCTTCCACAGCTGTCTTATCTC		1271
179	ThrPheIleProSerAspAspIleTyrAspCysLysValGluHisIstirpGlyLeuGlu		198
1272	ACCTTCATCCCTTCTGATGATGATTTATGACTGCAAGGTGGAGCAGCTGGGCGCTGGAG		1331
199	GluProValLeuLysHisIstirpGluProGluIleProAlaProMetSerGluLeuThrGlu		218

Db	1332	GAGCCGGTTCTGAACACCTGGGAACCTGAGATTTCAGCCCCCCTGAGAGCTGACAGNA	1391
Qy	219	Thr 219	
Db	1392	ACT 1394	
RESULT 14			
AAT60698			
ID	AAT60698	standard; cDNA; 588 BP.	
AC	AAT60698;		
XX			
DT	27-AUG-2003	(revised)	
DT	12-SEP-1997	(first entry)	
XX			
DE	Alpha2 region of Class II NOD mouse MHC (IAG7) cDNA.		
XX			
KW	Soluble; fusion; major histocompatibility complex; MHC; region;		
KW	heterodimer; complex; alpha2; antigen; binding groove; tolerance;		
KW	autoantigen; disease; insulin dependent; diabetes mellitus; IDDM;		
KW	antagonist; T cell; anergy; presenting cell; NOD mouse; Class II; alpha1;		
KW	ds.		
XX			
OS	Mus sp.		
XX			
PH	Key	Location/Qualifiers	
FT	mat_peptide	1..588	
XX		/*tag= a	
XX			
PN	WO9640944-A2.		
XX			
PD	19-DEC-1996.		
XX			
PF	07-JUN-1996;	96WO-US010102.	
XX			
PR	07-JUN-1995;	95US-00480002.	
PR	07-JUN-1995;	95US-00482133.	
PR	07-JUN-1995;	95US-00483241.	
PR	27-OCT-1995;	95US-0005964P.	
XX			
PA	(ZYMO) ZYMOGENETICS INC.		
PA	(ANER-) ANERGEN INC.		
XX			
PI	Kindsvogel W, Reich EP, Gross JA, Deshpande S, Sheppard PO;		
XX			
DR	WPI; 1997-052337/05.		
DR	P-PSDB; AAW10505.		
XX			
PT	Novel fused major histocompatibility complex:antigenic peptide complex -		
PT	useful to induce tolerance to an autoantigen-related disease e.g. insulin		
PT	-dependent diabetes mellitus.		
XX			
PS	Example 3; Page 132-133; 142pp; English.		
XX			
CC	A novel soluble fused major histocompatibility complex (MHC)		
CC	heterodimer:peptide complex, comprises DNA encoding 1st and 2nd MHC		
CC	domains, e.g. the present sequence, linked by DNA encoding a 5-25 residue		
CC	linker, and a DNA encoding an antigenic peptide able to associate with a		
CC	peptide binding groove of the MHC molecule, linked in frame to the DNA		
CC	encoding the 2nd domain by a DNA encoding a 5-25 residue linker. The		
CC	complex can be used to induce immunological tolerance in adults		
CC	susceptible to, or suffering from an autoantigen related disease, e.g.		
CC	insulin dependent diabetes mellitus (IDDM), by antagonising the binding		
CC	of particular T cells and antigen presenting cells, to induce anergy		
CC	(immunological non-responsiveness) in the targeted T cell. As the		
CC	heterodimers and corresponding antigen are permanently linked into a		
CC	single chain, obviating the requirement for complex heterodimer		
CC	truncation or formation, the complex eliminates inefficient and non-		
CC	specific peptide loading. (Updated on 27-AUG-2003 to correct OS field.)		
XX			
SQ	Sequence 588 BP; 153 A; 153 C; 136 G; 146 T; 0 U; 0 Other;		

Alignment Scores:
 Ad. No.: 3,51e-97 Length: 588
 Score: 1065.00 Matches: 196
 Percent Similarity: 100.0% Conservative: 0
 Best Local Similarity: 100.0% Mismatches: 0
 Query Match: 71.2% Indels: 0
 Gaps: 0

-10-048-116B-2_COPY_1_278 (1-278) x AAT60698 (1-588)

24 GluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrThrValTyrGlnSerPro 43
 |||||
 1 GAAGACGACATTGAGCGCCAGCACGATAGCTTCTATGTTACAACTGTTTATCATGCTCCT 60
 |||||
 44 GlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyrValAspLeu 63
 |||||
 61 GGAGACATTGGCCAGTACACATGAATTTGATGGTGTGATGGTGTCTATGTTGGACTTG 120
 |||||
 64 AspLysLysValThrValTyrPheGluProGluPheGlyGlnLeuLeuPheGluPro 83
 |||||
 121 GATAAGAGAAACTGCTGGAGGCTTCTGAGTTTGGCCAAATGTATCTTTGAGCCC 180
 |||||
 84 GlnGlyGlyLeuGlnAsnIleAlaGluLysHisAsnLeuGlyIleLeuThrLysArg 103
 |||||
 181 CAAAGTGGACTGCANAACATAGCTGCAGAAAACACAACTTGGGAATCTTGACTAAGAG 240
 |||||
 104 SerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPheProLysSerPro 123
 |||||
 241 TCAAAATTTTCAAGGCTACCAATGAGGCTCTCAAGCGACTGTGTTCCTCCCAAGTCCCT 300
 |||||
 124 ValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePheProVal 143
 |||||
 301 GTGCTGCTGGGTGAGCCCAACACCTTATCTGCTTTGTGGACAACATCTTCCACCTGTG 360
 |||||
 144 IleAsnIleThrTriPheuArgAsnSerLysSerValThrAspGlyValTyrGluThrSer 163
 |||||
 361 ATCAACATCACATGGCTCAGAAATAGCAAGTCAGTCACAGACGGCGTTTATGAGACGAG 420
 |||||
 164 PheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrIleuThrPheIleProSer 183
 |||||
 421 TTCTCTGTCACCGTGACCATTCCTTCCACAACTGCTTATCTCACCTTTCATCCCTTCT 480
 |||||
 184 AspAspAspIleTyrAspCysLysValGluHisTyrGlyLeuGluGluProValLeuLys 203
 |||||
 481 GATGATGACATTATGACTGCAGAGTGGAGCACTGGGGGCTTGAGAGCGGTTCTGAAA 540
 |||||
 204 HisTriPheProGluIleProAlaProMetSerGluLeuThrGluThr 219
 |||||
 541 CACTGGGAACCTGAGATTCCAGCCCCCATGTCTCAGAGCTGCAGAGACT 588
 |||||

3ULT 15
 235054

AAQ35054 standard; DNA; 776 BP.

AAQ35054;

25-MAR-2003 (revised)

21-MAY-1993 (first entry)

IAB alpha chain (known).

Acetylcholine receptor; epitope; myasthenia gravis; MG; antigen; binding site; MHC; IAB; alpha; beta; da.

Synthetic.

W09218150-A1.

29-OCT-1992.

23-APR-1992; 92WO-US0003391.

23-APR-1991; 91US-00690840.

PR 14-APR-1992; 92US-00869293.
 XX (ANER-) ANERGEN INC.
 XX Clark BR, Sharma SD, Lerch BL;
 XX WPI; 1993-036056/04.
 XX
 PT Pure major MHC-peptide complex - useful in treating deleterious immune
 PT response such as autoimmunity.
 XX Disclosure; Fig 8; 93pp; English.
 XX
 CC The ACHR peptide 195-215, which has been characterised as an epitope in
 CC myasthenia gravis (MG) in humans and in mice, may be connected to the N-
 CC terminal antigen binding site of a polypeptide derived from an MHC
 CC antigen associated with MG. For example, if the recombinant complex is to
 CC be used in mice, the ACHR peptide may be incorporated into a sequence
 CC encoding either the I-Ab-alpha or I-Ab-beta chain (see AAQ35054-55
 CC respectively). If the ACHR peptide is to be incorporated into the beta
 CC chain, for example, the oligonucleotide may be inserted as a replacement
 CC for the leader sequence. (Updated on 25-MAR-2003 to correct PN field.)
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX
 SQ Sequence 776 BP; 178 A; 211 C; 192 G; 194 T; 0 U; 1 Other;

Alignment Scores:
 Pred. No.: 1,04e-95 Length: 776
 Score: 1052.00 Matches: 195
 Percent Similarity: 93.2% Conservative: 9
 Best Local Similarity: 89.0% Mismatches: 15
 Query Match: 70.3% Indels: 0
 Gaps: 2

US-10-048-116B-2_COPY_1_278 (1-278) x AAQ35054 (1-776)

QY 1 MetProCysSerArgAlaLeuLeuLeuGlyValLeuAlaLeuAsnThrMetLeuSerLeu 20
 |||||
 Db 6 ATGCGCGCGAGGAGAGCTCTGATTTGGGGCTCTCGCCCTGACCACTGCTCACCCCTC 65
 |||||
 QY 21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrValTyr 40
 |||||
 Db 66 TGTGCGAGGTGAAGACGACATTGAGCGCCAGCAGTAGGCACCTATGTTAGTGTATAT 125
 |||||
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Ch completed: May 31, 2006, 23:18:27
time : 562.439 secs

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RESULT 2

BD137962

LOCUS

DEFINITION

BD137962 1446 bp DNA linear PAT 18-SEP-2002

Monovalent MHC-binding domain fused proteins and conjugates,

polyvalent MHC-binding domain fused proteins and conjugates,

polymer MHC-binding domain fused proteins and conjugates, and

utilization thereof.

ACCESSION

BD137962

VERSION

BD137962.1 GI:23232907

KEYWORDS

JP 2002504342-A/7.

SOURCE

synthetic construct

ORGANISM

other sequences; artificial sequences.

REFERENCE

1 (bases 1 to 1446)

Wucherpfennig,K.W. and Strominger,J.L.

Monovalent MHC-binding domain fused proteins and conjugates,

polyvalent MHC-binding domain fused proteins and conjugates,

polymer MHC-binding domain fused proteins and conjugates, and

utilization

JOURNAL

Patent: JP 2002504342-A 7 12-FEB-2002;

PRESIDENT AND FELLOWS OF HARVARD COLLEGE

OS Artificial Sequence

PN JP 2002504342-A/7

PD 12-FEB-2002

PF 19-FEB-1999 JP 2000532537

PR 19-FEB-1998 US 60/075351

PI KAI W WUCHERPFENNIG, JACK L STROMINGER

PC C12N15/09,A61K35/14,A61K47/48,C07K14/705,C07K16/00,C07K19/00,

PC C12Q1/02.

PC G01N33/53,C12N15/00

CC Description of Artificial Sequence: DR2-IgG fusion CC 3' end

of secretory signal

CC DRA*0101 extracellular domain

CC Linker

CC Pos leucine zipper domain

CC IgG domain

Key Location/Qualifiers

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FT misc_feature (730)..(1437).

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    AUTHORS
        Dreher, I. and Moll, T.
    TITLE
        Antagonists il-15
    JOURNAL
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        F. HOPFMANN-LA ROCHE AG (CH)
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DB	559	GTGAGCAGGATGACCCAGATGTCCAGATCAGCTGGTGTGTGAAACAACGTGGGAAGTACAC	618
QY	1000	ACAGCTCAGACACAAACCCCATAGAGAGGATTAACAACAGTACTCTCCGGTGGTCACTGCC	1059
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DEFINITION	Sequence 27 from Patent WO2004035622.		
ACCESSION	CQ806532		
VERSION	CQ806532.1	GI:47111926	
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SOURCE	synthetic construct		
ORGANISM	synthetic construct		
	other sequences; artificial sequences.		
REFERENCE	1		
AUTHORS	Dreher, I. and Moll, T.		
TITLE	Antagonists il-15		
JOURNAL	Patent: WO 2004035622-A 27 29-APR-2004;		
	F. HOFFMANN-LA ROCHE AG (CH)		

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ORIGIN		Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi; Muridea; Murinae; Mus.
Query Match	47.2%; Score 701.2; DB 2; Length 1108;	1 McLean,G.R., Nakouzi,A., Casadevall,A. and Green,N.S.
Best Local Similarity	93.3%; Pred. No. 1.3e-179;	2 Human and murine immunoglobulin expression vector cassettes
Matches	733; Conservative 0; Mismatches 53; Indels 0; Gaps 0;	3 Mol. Immunol. 37 (14), 837-845 (2000)
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		McLean,G.R.
		Direct Submission
		Submitted (18-SEP-2000) McLean G.R., Cell Biology, Albert Einstein
		College of Medicine, 1300 Morris Park Avenue, Bronx, New York
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Db	351	CTTGGGTGACCATCGTCTTCATCTTCCCTCCAAGATCAAGGATGATCATGATCTC 410
QY	906	CCTGAGCCCCATAGTCACATGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 965
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QY	1026	GGATTACACAGTCTCTCCGGGTGTGTCAGTGGTCCCTCCCATCCACGACGACCTGGAT 1085
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ION	V00798.1	GI:51835
ORDS	complementary DNA; gamma-immunoglobulin; immunoglobulin.	
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GANISM		
RENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
THORS	Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;	
TLE	Sciurognathi; Muridae; Murinae; Mus.	
URNAL	1. (bases 1 to 1095)	
UBMED	Sikorav,J.I., Aufray,C. and Rougeon,F.	
URES	Structure of the constant and 3' untranslated regions of the murine	
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IN		
ery Match	47.2%; Score 700.6; DB 6; Length 1095;	
st Local Similarity	100.0%; Pred. No. 1.9e-179;	
tches 700; Conservative	0; Mismatches 0; Indels 0; Gaps 0;	
786	TGAGCCCAAGGGGCCCAATCAAGCCCTGTCTCCATGCAAAATGCCAGACCTAACCT	845

291	TGAGCCCAAGGGGCCCAATCAAGCCCTGTCTCCATGCAAAATGCCAGACCTAACCT	350
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906	CCTGAGCCCCATAGTCACATGTGTGGTGGATGTGAGCGAGGATGACCCAGATGTCCA	965
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966	GATCAGCTGGTTTGTGAACAACAGTGAAGTACACACAGCTCAGACACAAACCCATAGAGA	1025
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1386	GAAGAACCTGGGTGGAAAGAAATAGCTACTCTCTGTTTTCAGTGGTTCACGAGGGTCTGCACAA	1445
891	GAAGAACCTGGGTGGAAAGAAATAGCTACTCTCTGTTTTCAGTGGTTCACGAGGGTCTGCACAA	950
1446	TCACACACAGCTAAGAGCTTCTCCGGACTCCGGGTAAA	1485
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DEFINITION	Sequence 4 from Patent EP 0338767.	
ACCESSION	I07390	
VERSION	I07390.1	GI:589918
KEYWORDS	Unknown.	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 1341)	
AUTHORS	Beavers,L.S.; Bumol,T.F.; Gadski,R.A. and Weigel,B.J.	
TITLE	Novel recombinant and chimeric antibodies directed against a human	
JOURNAL	adenocarcinoma antigen	
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Best Local Similarity	100.0%; Pred. No. 1.9e-179;	
Matches 700; Conservative	0; Mismatches 0; Indels 0; Gaps 0;	
QY	786	TGAGCCCAAGGGGCCCAATCAAGCCCTGTCTCCATGCAAAATGCCAGACCTAACCT 845

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RESULT 9
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DEFINITION
SEQUENCE 1 from Patent WO2005068503.
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WORDS
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ORGANISM
Mus musculus (house mouse)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muroidea; Muridae; Murinae; Mus.
1
Li.C.
REFERENCE
AUTHORS
TITLE
JOURNAL
PUBMED
12163274
REFERENCE 2 (bases 1 to 1407)
47.2%; Score 700.6; DB 2; Length 1401;

Best Local Similarity 100.0%; Pred. No. 1.9e-179;
Matches 700; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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AF466698 1407 bp mRNA linear ROD 17-DEC-2002
Mus musculus strain BALB/c immunoglobulin heavy chain mRNA, partial cds.
AF466698
AF466698.1 GI:27127159
ACCESSION
VERSION
KEYWORDS
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muroidea; Muridae; Murinae; Mus.
1 (bases 1 to 1407)
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AUTHORS
TITLE
JOURNAL
PUBMED
12163274
REFERENCE 2 (bases 1 to 1407)

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BD057272
LOCUS BD057272 1570 bp DNA linear PAT 29-SEP-1999
DEFINITION Gene encoding 6 from patent US 5859205.
ACCESSION AR029102
VERSION AR029102.1 GI:5941075
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 1570)
AUTHORS Adair, J. Robert., Athwal, D. Singh, and Emtage, J. Spencer.
TITLE Humanised antibodies
JOURNAL Patent: US 5859205-A 6 12-JAN-1999;
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Query Match 47.2%; Score 700.6; DB 2; Length 1570;
Best Local Similarity 100.0%; Pred. No. 2e-179;
Matches 700; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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BD057272
LOCUS BD057272 1570 bp DNA linear PAT 27-AUG-2002
DEFINITION Gene encoding antimalarial monoclonal antibody.
ACCESSION BD057272
VERSION BD057272.1 GI:22602878
KEYWORDS JP 2001275682-A/9.
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 1570)
AUTHORS Okawa, H., Nakata, M. and Yuasa, Y.
TITLE Gene encoding antimalarial monoclonal antibody
JOURNAL Patent: JP 2001275682-A 9 09-OCT-2001;
COMMENT KANKYO MENEKI GIJUTSU KENKYUSYO KK
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FH Key Location/Qualifiers.

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ORIGIN

Query Match 47.2%; Score 700.6; DB 2; Length 1570;
Best Local Similarity 100.0%; Pred. No. 2e-179;
Matches 700; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
786 TGAGCCACAGGGGCCACCAATCAAGCCCTGCTCTCCATGCAATGCCAGCACCTAACCT 845
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9372
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GANISM
RENC
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GBX;
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source
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st Local Similarity 100.0%; Pred. No. 2e-179;
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DB 1105 AACCATCTCAAAACCCAAAGGCTCAGTAAGAGCTCCACAGGTATATGTTTGCCTCCACC 1164
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DB 1165 AGAAGAAGAGATGACTAAGAAACAGGTCACTCTGACCTGCATGGTCAAGAGCTTCATGCC 1224
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QY 1326 TGAACCAAGTCTCGGACTCTGATGGTCTTCAATGATGACAGCAAGCTGAGAGTGGAAAA 1385
DB 1285 TGAACCAAGTCTCGGACTCTGATGGTCTTCAATGATGACAGCAAGCTGAGAGTGGAAAA 1344
QY 1386 GAAGAACTGGGTGGAAAGAAATAGTACTCTCGGACTCCGGGTAA 1445
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RESULT 15
AR559698 AR559698 1570 bp DNA linear PAT 08-OCT-2004
LOCUS Sequence 6 from patent US 6750325.
DEFINITION AR559698
ACCESSION AR559698
VERSION AR559698.1 GI:53969764
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 1570)
AUTHORS Jolliffe,L.K., Zivin,R.A., Adair,J.R. and Athwal,D.S.
TITLE CD3 specific recombinant antibody
PATENT: US 6750325-A 6 15-JUN-2004;
JOURNAL Celltech R&D Limited; Slough;
GBX;
FEATURES
source
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/organism="unknown"
/mol_type="genomic DNA"
ORIGIN
Query Match 47.2%; Score 700.6; DB 2; Length 1570;
Best Local Similarity 100.0%; Pred. No. 2e-179;
Matches 700; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
786 TGAGCCAGAGGCGCCACAAATCAAGCCCTGTCTCCATGCAAAATGCCAGCACCCTAACCT 845
745 TGAGCCAGAGGCGCCACAAATCAAGCCCTGTCTCCATGCAAAATGCCAGCACCCTAACCT 804
846 CTTGGTGGACCATCCGTCTTCATCTTCCCTCAAAGATCAAGGATGTAATCATGATCTC 905
805 CTTGGTGGACCATCCGTCTTCATCTTCCCTCAAAGATCAAGGATGTAATCATGATCTC 864
906 CTTGAGCCCATAGTACATGTTGGTGGATGTGAGCGGAGGATGACCCAGATGTCCA 965

865 CCTGAGCCCCATAGTCAATGTGTGGTGGATGTGAGCGAGGATGACCCAGATGTCCA 924
966 GATCAGCTGGTTTGTGAACAACGTGGAAAGTACACACAGCTCAGACACAAACCCATAGAGA 1025
925 GATCAGCTGGTTTGTGAACAACGTGGAAAGTACACACAGCTCAGACACAAACCCATAGAGA 984
1026 GGATTACACAGTACTCTCCGGGTGGTCACTGCGCTCCCTCCCATCCAGCACCAGGACTGGAT 1085
985 GGATTACACAGTACTCTCCGGGTGGTCACTGCGCTCCCTCCCATCCAGCACCAGGACTGGAT 1044
1086 GAGTGGCAAGGAGTTCAAAATGCAAGGTCAACAAACAAGACCTCCAGCGGCCCATCGAGAG 1145
1045 GAGTGGCAAGGAGTTCAAAATGCAAGGTCAACAAACAAGACCTCCAGCGGCCCATCGAGAG 1104
1146 AACCATCTCAAAACCCAAAGGGTCAGTAAGAGCTCCAAGGTATATGTCTTGCTCCACC 1205
1105 AACCATCTCAAAACCCAAAGGGTCAGTAAGAGCTCCAAGGTATATGTCTTGCTCCACC 1164
1206 AGAAGAAGAGATGACTAAGAAACAGGTCACTCTGACCTGCATGGTCACAGACTTCATGCC 1265
1165 AGAAGAAGAGATGACTAAGAAACAGGTCACTCTGACCTGCATGGTCACAGACTTCATGCC 1224
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1225 TGAAGACATTTACGTGGAGTGGACCAACACGGGAAACAGAGCTAACTACAAGAACAC 1284
1326 TGAACCAAGTCTCGACTCTGATGGTCTTACTTCATGTACAGCAAGCTGAGAGTGGAAAA 1385
1285 TGAACCAAGTCTCGACTCTGATGGTCTTACTTCATGTACAGCAAGCTGAGAGTGGAAAA 1344
1386 GAAGAAGTGGTGGAAAGAAATAGTACTCTGTTCACTGTTCCAGGAGGCTTCGCACAA 1445
1345 GAAGAAGTGGTGGAAAGAAATAGTACTCTGTTCACTGTTCCAGGAGGCTTCGCACAA 1404
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1405 TCACCACACGACTAAGAGCTTCTCCCGGACTCCCGGGTAAA 1444

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nucleic - nucleic search, using sw model

on: May 31, 2006, 23:41:05 ; Search time 1026.58 Seconds
 (without alignments)
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ring table: IDENTITY_NUC
 Gapop 10.0 , Gapext 1.0

atched: 5244920 seqs, 3486124231 residues

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imum DB seq length: 2000000000

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 Maximum Match 100%
 Listing first 45 summaries

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- 2: geneseqn1990s:*
- 3: geneseqn2000s:*
- 4: geneseqn2001as:*
- 5: geneseqn2001bs:*
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- 11: geneseqn2003ds:*
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- 13: geneseqn2004bs:*
- 14: geneseqn2005s:*
- 15: geneseqn2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Full No.	Score	Query Match	Length	ID	Description
1	1484	99.9	1484	5	Aaf55098 DNA encod
2	1067.4	71.9	1676	4	Abi99041 Murine PC
3	957.6	64.5	1446	2	Aat99707 DR2-IgG F
4	957.6	64.5	1446	2	Aax87813 HLA-DR2 a
5	957.6	64.5	1446	14	Adw44282 DR2-IgG F
6	726.2	48.9	2346	4	Abi99027 IAS MBP 1
7	723.2	48.7	2343	4	Abi99033 MBP 90-10
8	701.2	47.2	1045	12	Ado07566 Fusion pr
9	701.2	47.2	1108	12	Ado07578 Fusion pr
10	701.2	47.2	1108	12	Ado07577 Fusion pr
11	700.6	47.2	990	12	Adl15694 Murine im
12	700.6	47.2	1401	14	Aec20762 M-CSF spe
13	700.6	47.2	1560	14	Aed19725 Anti-PpP
14	700.6	47.2	1569	14	Adv26108 Mouse OKT
15	700.6	47.2	1569	14	Adw71834 Murine OK
16	700.6	47.2	1570	12	Aaq12637 Monoclonal
17	700.6	47.2	1570	12	Adg91058 Murine OK
18	700.6	47.2	3973	13	Adt77690 Monoclonal

19	700.4	47.2	1509	14	AEE21942	Aee21942 Single ch
20	700.2	47.2	729	3	AAZ35704	Aaz35704 Human gly
21	700.2	47.2	1140	10	ADe85817	Ad85817 Murine in
22	700	47.1	1158	2	AAT59350	Aat59350 1-153 del
23	700	47.1	1188	2	AAT59349	Aat59349 1-153 c-m
24	700	47.1	6729	4	AAF30341	Aaf30341 Bicistron
25	700	47.1	7528	4	AAF30316	Aaf30316 Bicistron
26	699.4	47.1	1530	14	ABE12356	Aeb12356 Fusion pr
27	699.4	47.1	1530	14	AED64236	Aed64236 mFc-hOgH-
28	699	47.1	699	15	AEF05391	Aef05391 Human mFc
29	699	47.1	708	14	ABE12344	Aeb12344 Murine im
30	699	47.1	708	14	AED64224	Aed64224 Murine im
31	699	47.1	1341	1	AA91659	Aan91659 Chimeric
32	699	47.1	1413	14	ABE12360	Aeb12360 Fusion pr
33	699	47.1	1413	14	AED64240	Aed64240 hTSH-mFc-
34	699	47.1	1431	14	ABE12362	Aeb12362 Fusion pr
35	699	47.1	1431	14	AED64242	Aed64242 hOgH-mFc-
36	699	47.1	1581	2	AAQ48037	Aaq48037 Monoclonal
37	699	47.1	1645	2	AAQ54652	Aaq54652 T84.12 He
38	698.4	47.0	1131	2	AAV55416	Aav55416 Chimeric
39	698.4	47.0	1194	2	AAV55415	Aav55415 Chimeric
40	698.4	47.0	1275	2	AAT62850	Aat62850 Mouse sol
41	698.4	47.0	1473	13	ADS31748	Ad31748 DNA encod
42	698.4	47.0	1473	13	ADS92750	Ad92750 DNA encod
43	697.8	47.0	1524	14	ABE21946	Aee21946 Single ch
44	697.4	47.0	699	3	AAZ51300	Aaz51300 Murine im
45	697.4	47.0	699	3	AAZ50055	Aaz50055 Mouse imm

ALIGNMENTS

RESULT 1
 AAF5098
 ID AAF5098 standard; DNA; 1484 BP.

AC AAF5098;
 DT 15-MAY-2001 (first entry)

DE DNA encoding a fusion protein comprising an alpha chain of MHC.

KW Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
 KW major histocompatibility complex; FC region; antigen; T lymphocyte;
 KW immunostimulant; vaccine; infection; tumour; ss.

OS Synthetic.

FH Key Location/Qualifiers
 CDS 1..1482
 FT /*tag= a

PN WO200109194-A1.

PD 08-FEB-2001.

PF 28-JUL-2000; 2000WO-FR002193.

PR 29-JUL-1999; 99FR-00009862.

PA (CNRS) CNRS CENT NAT RECH SCI.

PI Glaichenhaus N, Malherbe L;

XX WPI; 2001-182944/18.

DR P-PSDB; AAB67480.

PT New soluble recombinant protein, useful e.g. as immunostimulant,
 PT comprises dimeric major histocompatibility complex molecule fused to
 PT immunoglobulin FC region.

PS Example 1; Page 31-33; 43pp; French.

XX

22-MAR-2001; 2001WO-US009616.

22-MAR-2000; 2000US-0191274P.

15-MAY-2000; 2000US-0204249P.

23-JAN-2001; 2001US-0264003P.

(CORI-) CORIXA CORP.

Carter D, Zhu S, Arimilli S, Wang A;

WPI; 2001-616371/71.

P-PSDB; ABB56471.

Multimeric complex for treating autoimmune diseases, comprises first and second single chain MHC class II molecules, each comprising alpha1 and beta1 domain linked through amino acid linker and multimerization domain.

Disclosure; Page 115; 147pp; English.

The invention relates to a multimeric complex comprising a first recombinant single chain major histocompatibility complex (MHC) class II molecule and a second recombinant single chain MHC class II molecule, each comprising an alpha1 domain and a beta1 domain linked through an amino acid linker and a multimerization domain. The first and the second molecule are linked through the multimerization domain to form a multimeric complex. The complex is useful for treating autoimmune diseases. It is useful for treating insulin dependent diabetes, multiple sclerosis, myasthenia gravis, pernicious anaemia, autoimmune encephalomyelitis (EAE), rheumatoid arthritis and systemic lupus erythematosus. The present sequence encodes a single chain MHC class II molecule of the invention

Sequence 1676 BP; 438 A; 470 C; 407 G; 361 T; 0 U; 0 Other;

Query Match 71.9%; Score 1067.4; DB 4; Length 1676;
 Fast Local Similarity 83.5%; Pred. No. 2.9e-277;
 Matches 1319; Conservative 0; Mismatches 96; Indels 164; Gaps 3;

67 GGTGAAGACGACATTGAGCGCCGACACGATGGCTTCTATGGTACAACTGTTATCAGTCT 126

91 GGCGAAGACGACATTGAGCGCCGACACGATGGCTTCTATGGTACAACTGTTATCAGTCT 150

127 CCTGGAGACATTGGCCAGTACACACATGAATTTGATGGTGATGATGTTCTATGGGAC 186

151 CCTGGAGACATTGGCCAGTACACACATGAATTTGATGGTGATGATGTTCTATGGGAC 210

187 TTGGATAAGAGAAAACCTGTCTGGAGCTTCTCTGAGTTTGGCCAAATTTGATCTCTTTGAG 246

211 TTGGATAAGAGAGAACTATCTGGATGCTTCTGAGTTTGGCCAAATTTGATCTCTTTGAC 270

247 CCCCAAGGTGGACTGCAAAACATAGCTGCGAGAAAAACAACATTTGGAAATCTTGACTAAG 306

271 CCCCAAGGTGGACTGCAAAACATAGCTGCGAGAAAAACAACATTTGGAAATCTTGACTAAG 330

307 AGGTCAAAATTCACCCAGCTACCAATGAGGCTCTCAAGCGACTGTGTCCCAAGTCC 366

331 AGGTCAAAATTCACCCAGCTACCAATGAGGCTCTCAAGCGACTGTGTCCCAAGTCC 390

367 CCTGTGCTGTGGGTGAGCCCAACACCTTATCTGTCTTGTGGCAACAATCTTCCCACT 426

391 CCTGTGCTGTGGGTGAGCCCAACACCTTATCTGTCTTGTGGCAACAATCTTCCCTCT 450

427 GTGATCAACATCAGTGGCTCAGAAATAGCAAGTCAAGTCAAGCGCGTTATAGAGCC 486

451 GTGATCAACATCAGTGGCTCAGAAATAGTAACTCAAGTCAAGCGCGTTATAGAGCC 510

487 AGCTTCTGTGTAACCGTGAACATTCCTTCCCAAGCTGTCTTATCTCACTTATCCCT 546

511 AGCTTCTGTGTAACCGTGAACATTCCTTCCCAAGCTGTCTTATCTCACTTATCCCT 570

547 TCTGATGATGACATTTATGACTGCAAGGTGGACACTGGGGCTGGAGGAGCGGTTCTG 606

Db	571	TCCTGACGATGATATTTATGACTGCAAGGTGGAGCACTGGGGCTCGGAGGCGGTTCTG	630
Qy	607	AAACACTGGGAACCTGAGATTTCCAGCCCCCATGTGAGAGCTGACAGAAACTGGAGTGA	666
Db	631	AAACACTGGGAACCTGAGATTTCCAGCCCCCATGTGAGAGGATCTGCCAAAACAAGCC	690
Qy	667	GGAT-----	670
Db	691	CCATCGGTCTATCCACTGGCCCTGTGTGGGAGATACAACCTGGCTCCTCGGTGACTCTA	750
Qy	671	-----	670
Db	751	GGATCGGTCTGATCAAGGGTTATTTCCCTGAGCCAGTGACCTTGACCTGGAACCTGGATCC	810
Qy	671	-----	686
Db	811	CTGTCCAGTGGTGTGCACACTTCCAGCTGTCTGTCAGTCTGACCTCTACACCTCAGC	870
Qy	687	AGCTCAGCTCGAAAAAAGAGCTCCAGGCCCTGGAGAAAGGAAAATGCAAGCTGGAATGGGA	746
Db	871	AGCTCAG--TGACTGTAACCTCGAGCACTGGCCAGCCAGTCCATCACTGCAATGTGG	928
Qy	747	GTTCGAAGCACTGGAAAAAGAACTGGCTCAGGCAAGCATCTGAGCCAGAGGGCCCAAT	806
Db	929	CCACCCGGCAAGCAGCAGCAAGGTGGAAGAA--AATTGAGCCAGAGGGCCCAAT	986
Qy	807	CAAGCCCTGTCTCCATGCAAAATGCCAGCACTTAACCTCTTGGGTGGACCATCCCTCT	866
Db	987	CAAGCCCTGTCTCCATGCAAAATGCCAGCACTTAACCTCTTGGGTGGACCATCCCTCT	1046
Qy	867	CATCTTCCCTCCAAAGATCAAGGATGTACTCATGATCTCCCTGAGCCCATAGTCAATG	926
Db	1047	CATCTTCCCTCCAAAGATCAAGGATGTACTCATGATCTCCCTGAGCCCATAGTCAATG	1106
Qy	927	TGTGTGGTGGATGTGAGCGAGATGACCCAGATGTCCAGATCAGCTGGTTTGTGAACAA	986
Db	1107	TGTGTGGTGGATGTGAGCGAGATGACCCAGATGTCCAGATCAGCTGGTTTGTGAACAA	1166
Qy	987	CGTGAAGTACACACAGCTCAGACACAAACCCATAGAGAGGATTACAAAGTACTCTCCG	1046
Db	1167	CGTGAAGTACACACAGCTCAGACACAAACCCATAGAGAGGATTACAAAGTACTCTCCG	1226
Qy	1047	GGTGTGTCAGTGCCTCCCATCCAGCAGCAGCTGGATGAGTGAGGCAAGAGTTCAAAATG	1106
Db	1227	GGTGTGTCAGTGCCTCCCATCCAGCAGCAGCTGGATGAGTGAGGCAAGAGTTCAAAATG	1286
Qy	1107	CAAGGTCAAACAAGAGCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCCAAGG	1166
Db	1287	CAAGGTCAAACAAGAGCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCCAAGG	1346
Qy	1167	GTGATGAGCTCCACAGGTATATGTCTTGGCTCCACAGAGAGAGATGACTAAGAA	1226
Db	1347	GTGATGAGCTCCACAGGTATATGTCTTGGCTCCACAGAGAGAGATGACTAAGAA	1406
Qy	1227	ACAGGTCACTCTGACCTGATGTGTCAAGACTTTCATGCTGAGAGACATTTACGTGAGTG	1286
Db	1407	ACAGGTCACTCTGACCTGATGTGTCAAGACTTTCATGCTGAGAGACATTTACGTGAGTG	1466
Qy	1287	GACCAACAACGGGAAAAACAGAGCTAAACTACAAGAACACTGAACCACTGAGCTCTGA	1346
Db	1467	GACCAACAACGGGAAAAACAGAGCTAAACTACAAGAACACTGAACCACTGAGCTCTGA	1526
Qy	1347	TGGTTCTTACTTATGATGATGAGCAAGCTGGAAGAGAGAGATGAGTGGAGAGAA	1406
Db	1527	TGGTTCTTACTTATGATGATGAGCAAGCTGGAAGAGAGAGATGAGTGGAGAGAA	1586
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Db	1587	TAGTCTCTCTGTTCAGTGGTCCAGAGGCTTGCAACATCTGCAACATCACCACCTAAGAGCTT	1646
Qy	1467	CTCCCGGACTCCGGGTAAA 1485	
Db	1647	CTCCCGGACTCCGGGTAAA 1665	

..LT 3

ART99707 standard; cDNA; 1446 BP.

AAT99707;

17-OCT-2003 (revised)

17-AUG-1998 (first entry)

DR2-IgG fusion construct.

Major histocompatibility complex class II; MHC class II; human; mouse; fusion protein; HLA-DR2; DRA*0101; binding domain; Fos; dimerisation domain; IgG; allergy; autoimmune disease; vaccine; multiple sclerosis; therapy; ss.

Homo sapiens.

Mus musculus.

Chimeric.

WO9806749-A2.

19-FEB-1998.

15-AUG-1997; 97WO-US014503.

16-AUG-1996; 96US-0024077P.

(HARD) HARVARD COLLEGE.

Wucherpennig KW, Strominger JL;

WPI; 1998-159459/14.

New Class II MHC fusion proteins - comprising a MHC Class II binding domain and a dimerisation domain or an immunoglobulin region used for modulating immune responses.

Example; Page 49; 76pp; English.

This nucleotide sequences codes for a bivalent DR2 fusion protein obtained by fusion of the Fc portion of IgG2a to the 3' end of a DR-alpha-Fos cDNA construct (see AA16866). The Fc portion was amplified by RT-PCR from mouse hybridoma L243. The PCR product was then fused in frame with the DR-alpha-Fos construct by overlapping PCR. The DR2-IgG fusion was expressed in the prosofilla Schneider cell system. The invention relates to new soluble monovalent and multivalent Class II MHC fusion proteins comprising a MHC Class II binding domain and a dimerisation domain or an immunoglobulin region that can be used for the treatment of allergic and autoimmune diseases (e.g. multiple sclerosis), for tolerising a subject to foreign tissue before or after organ or tissue transplantation, or for vaccination against pathogens. (Updated on 17-OCT-2003 to standardise OS field)

Sequence 1446 BP; 414 A; 375 C; 356 G; 301 T; 0 U; 0 Other;

ery Match 64.5%; Score 957.6; DB 2; Length 1446;

et Local Similarity 81.8%; Pred. No. 1.2e-247;

tches 1136; Conservative 0; Mismatches 234; Indels 18; Gaps 2;

116 TTTATCAGTCTCTGGAGACATTTGCCAGTACACATCAATTTTCATGGTGTATGATTTGT 175

50 TCTATCTGATCTCTGACCATCAGGGAGTTTATGTTGACTTTCATGGTGTATGATTTT 109

176 TCTATGTGGACTTGGATAGAGAAACCTGCTCGAGGCTTCTCGAGTTTGGCCAAATTGA 235

110 TCCATGTGGATATGGCAAGAGGAGACGGTCTCGGGCTTGAAGAAATTTGGACGATTTG 169

236 TACTCTTTGGCCCCCAAGGTGACATGCGAAACATAGCTGCGAAACACACACTTGGGA 295

170 CCAGCTTTTGAGGCTCAAGTGCATTTGGCCCAACATAGCTGTGGACAAAGCCAACTTGGAAA 229

QY 296 TCTTGACTAAGAGGTCAAATTTTCACTCCAGCTACCAATAGAGGCTCTCTCAAGCGACTGTGT 355
 DB 230 TCATGACAAAGCGTCCAACTATATCTCGATCACCAATGTACCTCCAGAGGTAACTGTGC 289
 QY 356 TCCCAAGTCCCTGTGTCTGGTTCAGCCCAACACCTTATCTGCTTTGTGGACAACA 415
 DB 290 TCACGAACAGCCCTGTGGAACTGAGAGAGAGCCCACTCTCATCTGTTTCATAGACAGT 349
 QY 416 TCTTCCACCTGTGATCAACATCACTGGCTCAGAAATAGCAAGTCAGTCACAGACGGGG 475
 DB 350 TCACCCACCAAGTGTCAATGTACGTGGCTTCGAAATGGAJAACTGTCCACACAGAG 409
 QY 476 TTTATGAGACAGAGTTCCTCGTCAACCGTGACCAATTCCTTCCAAGCTGTCTTATCTCA 535
 DB 410 TGTGAGACAGAGTCTTCTGCTGCCAGGGAAGACCACTTTTCCGCAAGTTCACATATCTCC 469
 QY 536 CTTTCATCCCTTCTGATGATGACATTTATGACTGCAAGGTGGAGCACTGGGGCTTGGAGG 595
 DB 470 CTTTCTGCTCCCTCAACTGAGGACGTTTACGACTGCAGGGTGGAGCACTGGGGCTTGGATG 529
 QY 596 AGCGGTTCTGAJAAACACTGGGAACCTGAGATTTCAGCCCCCATGTCCAGAGCTGACAGAAA 655
 DB 530 AGCTCTTCTCAAGCACTGGAGATTGATGCTCCAGCCCTCTCCAGAGACTACAGAGG 589
 QY 656 CTGGAGGTGGAGGATCCACT-----ACAGCTCCATCAGCTCAGCTCGAAAA 700
 DB 590 TCGACGGAGGTGGCGGGGTTTAACTGATACACTCCAAGCGAGAGACAGATCAACTTGAAG 649
 QY 701 AAGAGCTCCAGGCCCTGGAGAAAGGAATGACAGCTGGATGGAGTTGCAAGCACTGG 760
 DB 650 ACAGAGAGTCTGCGTTGCAGACCGAGATTGCCAATCTACTGAAGAGAGGAAAAAAGTGG 709
 QY 761 AAAAGGAAGTCTGCTCAGGAGCATCTGAGCCAGAGGGCCCAATCAAGCCCTGTC 817
 DB 710 AGTTTCATCTGGCGCCCATCGAGATCTGAGCCAGAGGGCCCAATCAAGCCCTGTC 769
 QY 818 TCCATGCAAAATGCCCCAGCACCTAACTCTTTGGGTGGACCATCCCTCTTCATCTTCCCTC 877
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 QY 878 CAAAGATCAAGAGTGTACTCATGATCTCCCTGAGCCCATAGTACATGTGTGGTGGTG 937
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 QY 938 ATGTGAGCGAGATGACCCAGATGTCAGATCAGCTGCTTTGTGAACCAACCTGGAAGTAC 997
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 DB 1010 CCCTCCCATCCAGCACAGAGCTGGATGAGTGGCAAGAGTTTCAAAATGCAAGGTCAACA 1069
 QY 1118 CAAAGAGCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTCAGTAAGAG 1177
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 QY 1178 TCCACAGGTTATGTCTTGGCTCCACAGAGAGAGATGACTTAAGAAAAAGGTCACTC 1237
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 DB 1190 TGACCTGATGGTTCAGAGCTTCATGCTGAAGACATTTAGTGGAGTGGACCAACAG 1249
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 DB 1250 GGAJAAACAGAGCTAAACTACAAAGAACACTGAACCCAGTCTCTGAGTCTCTGATGGTCTTACT 1309


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770 CTCATGCAAAATGCCAGACCTAACTCTTGGTGGACCATCGCTCTTCATCTCCCTC 829
878 CAAGATCAAGATGATCTCATGATCTCCCTGAGCCCATAGTCACATGCTGGTGG 937
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830 CAAGATCAAGATGATCTCATGATCTCCCTGAGCCCATAGTCACATGCTGGTGG 889
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998 ACACAGCTCAGACACAAACCCATAGAGAGGATTACACAGTACTCTCCGGGTGTCAGTG 1057
950 ACACAGCTCAGACACAAACCCATAGAGAGGATTACACAGTACTCTCCGGGTGTCAGTG 1009
1058 CCCTCCCATCCAGCACGAGCTGGATGAGTGGCAAGGATTCAAAATGCAAGGTCACACA 1117
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1118 ACAAGAGCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCCAAGGGTCAGTAAGAG 1177
1070 ACAAGAGCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCCAAGGGTCAGTAAGAG 1129
1178 CTCACAGGTATATCTTGGCTTCCACAGAGAGAGATGACTAAGAAAACAGTCACTC 1237
1130 CTCACAGGTATATCTTGGCTTCCACAGAGAGAGATGACTAAGAAAACAGTCACTC 1189
1238 TGACCTGATGGTCAAGACTTCACTGCTGAGACATTTAGTGGAGTGGACCAACACG 1297
1190 TGACCTGATGGTCAAGACTTCACTGCTGAGACATTTAGTGGAGTGGACCAACACG 1249
1298 GGAACACAGAGCTAAACTACAAGAACTGAAACCACTCTGAGCTCTGATGGTTCTTACT 1357
1250 GGAACACAGAGCTAAACTACAAGAACTGAAACCACTCTGAGCTCTGATGGTTCTTACT 1309
1358 TCATGTCAGCAAGCTGAGAGTGGAAAAGAGAACTGGGTGGAAGAAATAGTACTCCT 1417
1310 TCATGTCAGCAAGCTGAGAGTGGAAAAGAGAACTGGGTGGAAGAAATAGTACTCCT 1369
1418 GTTCAGTGGTCCAGAGGCTGCAACATCACCACAGCTAAGAGCTTCTCCCGGACTC 1477
1370 GTTCAGTGGTCCAGAGGCTGCAACATCACCACAGCTAAGAGCTTCTCCCGGACTC 1429
1478 CGGTAAAA 1485
1430 CGGTAAAA 1437
24-MAR-2005 (first entry)
DR2-IgG fusion protein encoding DNA.
```

Major histocompatibility complex; fusion protein; immunoconjugate; adoptive immunotherapy; dermatological; immunosuppressive; antirheumatic; antiarthritic; neuroprotective; antiinflammatory; autoimmune diseases; pemphigus vulgaris; rheumatoid arthritis; multiple sclerosis; systemic lupus erythematosus; immune disorder; DR2-IgG protein; gene; ds.

Homo sapiens.
Chimeric.
Unidentified.

Key Location/Qualifiers
CDS 1..1440
/tag= b
/product= "DR2-IgG fusion protein"
/partial

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FT misc_feature /note= "No start codon"
FT 1..15 /tag= a
FT /note= "3' end of secretory signal"
FT 16..598 C
FT /tag= C
FT /note= "DRA*010extracellular domain"
FT 589..609
FT /tag= d
FT /note= "Linker sequence"
FT 610..729
FT /tag= e
FT /note= "Fos Leucine zipper domain"
FT 730..1437
FT /tag= f
FT /note= "IgG domain"
FT
XX
XX US2005003431-A1.
XX
XX 06-JAN-2005.
XX
XX 21-JUL-2004; 2004US-00895543.
XX
XX 16-AUG-1996; 96US-0024077P.
XX 15-AUG-1997; 97WO-US014503.
XX 19-FEB-1998; 98US-0075351P.
XX 12-FEB-1999; 99US-00248964.
XX
XX (WUCH/) WUCHERPENNIG K W.
XX (STRO/) STROMINGER J L.
XX
XX Wucherpennig KW, Strominger JL;
XX WPI; 2005-089945/10.
XX P-PSDB; ADM44283.
XX
XX Novel class II major histocompatibility complex (MHC) fusion protein
XX having MHC class II binding domain of MHC class II alpha chain, and
XX dimerization domain, useful for treating pemphigus vulgaris, rheumatoid
XX arthritis.
XX
XX Example; SEQ ID NO 11; 55pp; English.
XX
XX The present invention relates to the class II major histocompatibility
XX complex (MHC) fusion protein having MHC class II binding domain of MHC
XX class II alpha chain and a dimerization domain. The invention is useful
XX in adoptive immunotherapy and tolerizing against foreign tissue. The
XX invention is also useful for treating autoimmune diseases such as
XX pemphigus vulgaris, rheumatoid arthritis, multiple sclerosis and systemic
XX lupus erythematosus. The present sequence is the DR2-IgG fusion protein
XX encoding DNA.
XX
XX Sequence 1446 BP; 414 A; 375 C; 356 G; 301 T; 0 U; 0 Other;
XX
XX Query Match 64.5%; Score 957.6; DB 14; Length 1446;
XX Best Local Similarity 81.8%; Pred. No. 1.2e-247;
XX Matches 1136; Conservative 0; Mismatches 234; Indels 18; Gaps 2;
Qy 116 TTTATCATGCTCCCTGGAGACATTCGCCAGTACACACATGAATTTGATGGTGTGATGTTGT 175
Db 50 TCTATCTGAATCTCTGACCAATCAGCGGAGTTTATGTTGACTTTGATGGTGTGATGATTT 109
Qy 176 TCTATGTGGACTTCGATAAGAAAGAACTGCTCGAGGCTTCTCGAGTTTGGCCAAATGGA 235
Db 110 TCCATGTGGATATGCAAGAGAGAGGAGCGCTTCGCCGCTTGAAGAATTTGGACGATTG 169
Qy 236 TACTCTTTGAGCCCCAAGGTGGAATCGCAAAACATAGCTGCGAAGAAAACACAACTTGGGAA 295
Db 170 CCAGCTTTGAGGCTCAAGGTGCAATTTGGCCACATAGCTGTGGACAAAGCCAACTTGGAAA 229
Qy 296 TCTTGACTTAAGAGGTCAAAATTTTACCCTCAGCTACCAATGAGGCTCTCTCAAGGACTGTGT 355
Db 230 TCATGACAAAGCGCTCCAACTATATCTCCGATCACAATGTACTTCCAGAGGTAATGTGTC 289
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356 TCCCCAAGTCCCTGTGCTGCTGGGTGAGCCCAACACCCCTTATCTGCTTGTGGACAACA 415
290 TCACGAACAGCCCTGTGGAACTGAGAGAGCCCAACGCTCTCATCTGTTTCATAGACAAGT 349
416 TCTTCCACACCTGTGATCAACATCACATGCTGCTCAGAAATAGCAAGTCAGTCACAGAGCGGCG 475
350 TCACCCCAACAGTGTCAATGTCACTGCTGCTCGAAATGGAACCTGTTCACCCACAGGAG 409
476 TTTATGAGACCAAGTCTCTGCTCAACCGTGACATTCCTTTCACAAGCTGTCTTATCTCA 535
410 TGTTCAGAGACAGTCTCTCTGCCAGGGAAGACCACTTTTCCGCAAGTTCACATATCTCC 469
536 CCTTCACTCCCTCTGTATGATGACATTTATGACTGCAAGGTGGAGCACTGGGGCTCGAGG 595
470 CCTTCTGCTCCCTCAACTGAGAGGTTTACGACTGCAAGGTGGAGCACTGGGGCTTGGATG 529
596 AGCCGGTCTTGAACCACTGGGAACCTGAGATTCAGAGCCCTCATGTTCAGAGCTCAACAGAA 655
530 AGCTCTTCTCAAGCACTGGAGTGTGATGCTCCAGGCCCTCTCCAGAGACTACAGAGG 589
656 CTGGAGGTGAGGATCCACT-----ACAGCTCCATCAGCTCAGCTCGAAA 700
590 TCGACGAGGTGGCGCGGTAACTGATACACTCCAAAGCGGAGACAGATCAACTTGAAG 649
701 AAGAGCTCCAGGCCCTGGGAGAGGAATGCAAGCTGGATGGGAGTTGCAAGCACTGG 760
650 ACGAGAGTCTGCGTTGCAAGACCGAGATTTGCCAATCTACTGAAAGAGAGGAAGAACTGG 709
761 AAAAGGAACGTG---GCTCAGGAGAGCATCTGAGGCCAGAGGGGCCCAATCAAGCCCTGTC 817
710 AGTTCACTCTGGCGGCCCATGAGCATCTGAGGCCAGAGGGGCCCAATCAAGCCCTGTC 769
818 CTCCATGCAAAATGCCAGACACTTAACCTCTTGGGTGGAACATCCGCTTTCATCTTCCCTC 877
770 CTCATGCAAAATGCCAGACACTTAACCTCTTGGGTGGAACATCCGCTTTCATCTTCCCTC 829
878 CAAGATCAAGGTGACTCATGATCTCCCTGAGGCCCATAGTCAATGCTGTGGTGGTGG 937
830 CAAAGATCAAGGTGACTCATGATCTCCCTGAGGCCCATAGTCAATGCTGTGGTGGTGG 889
938 ATGTGAGCGAGGATGACCCAGATGTCCAGATCAGCTGCTGTTTGTGAACCAACGTCGAAGTAC 997
890 ATGTGAGCGAGGATGACCCAGATGTCCAGATCAGCTGCTGTTTGTGAACCAACGTCGAAGTAC 949
998 ACAAGCTCAGACACAAACCCATAGAGAGGATTAACAAGTACTCTCCGGGTGGTCAGTG 1057
950 ACACAGCTCAGACACAAACCCATAGAGAGGATTAACAAGTACTCTCCGGGTGGTCAGTG 1009
1058 CCTCCCTCCATCCAGCACAGGACTGGATGATGGTGGCAAGGATTCAAATGCAAGGTCAACA 1117
1010 CCTCCCTCCATCCAGCACAGGACTGGATGATGGTGGCAAGGATTCAAATGCAAGGTCAACA 1069
1118 ACAAGACCTCCAGCGGCCCATCGAGAGAACCATCTCAAAACCCAAAGGTCAGTAAGAG 1177
1070 ACAAGACCTCCAGCGGCCCATCGAGAGAACCATCTCAAAACCCAAAGGTCAGTAAGAG 1129
1178 CTCACAGGTATATGTCTTGGCTCCACAGAAAGAGATGATTAAGAAACAGGTCACTC 1237
1130 CTCACAGGTATATGTCTTGGCTCCACAGAAAGAGATGATTAAGAAACAGGTCACTC 1189
1238 TGACCTGCACTGTCACAGACTTCATGCTGGAAGACATTTACGTGGAGTGGCAACCAACG 1297
1190 TGACCTGCACTGTCACAGACTTCATGCTGGAAGACATTTACGTGGAGTGGCAACCAACG 1249
1298 GGAAGACAGAGCTAAACTACAAGAACACTGAACAGTCTCGGACTCTGTGTTCTTACT 1357
1250 GGAAGACAGAGCTAAACTACAAGAACACTGAACAGTCTCGGACTCTGTGTTCTTACT 1309
1358 TCATGTACAGCAAGCTGAGAGTGGAAAGAAAGAACTGGGTGGAAAGAAATAGTACTCTCT 1417
1310 TCATGTACAGCAAGCTGAGAGTGGAAAGAAAGAACTGGGTGGAAAGAAATAGTACTCTCT 1369

QY 1418 GTTCAGTGTCCACGAGGGTCTGCACAATCACCACACGACTAAGAGCTTCTCCGGACTC 1477
Db 1370 GTTCAGTGTCCACGAGGGTCTGCACAATCACCACACGACTAAGAGCTTCTCCGGACTC 1429
QY 1478 CGGGTAAA 1485
Db 1430 CGGGTAAA 1437
RESULT 6
ABI99027
ID ABI99027 standard; cDNA; 2346 BP.
XX
AC ABI99027;
XX
DT 25-FEB-2002 (first entry)
XX
TAS MBP 1-14 CH1.CH2.CH3 coding sequence.
XX
KW Mouse; MHC; major histocompatibility complex; MHC class II; multimer;
single chain; immunosuppressive; antidiabetic; antiinflammatory;
antianemic; antirheumatoid; antiarthritic; neuroprotective; vaccine;
autoimmune disease; insulin dependent diabetes; multiple sclerosis;
myasthenia gravis; pernicious anaemia; autoimmune encephalomyelitis;
rheumatoid arthritis; systemic lupus erythematosus; ss.
XX
OS Mus sp.
OS Synthetic.
XX
PN W0200170245-A1.
XX
PD 27-SEP-2001.
XX
PF 22-MAR-2001; 2001WO-US009616.
XX
PR 22-MAR-2000; 2000US-0191274P.
PR 15-MAY-2000; 2000US-0204249P.
PR 23-JAN-2001; 2001US-0264003P.
XX
PA (CORI-) CORIXA CORP.
XX
PI Carter D, Zhu S, Arimilli S, Wang A;
XX
WI: 2001-616371/71.
DR P-PSDB; ABB56457.
XX
PT Multimeric complex for treating autoimmune diseases, comprises first and
second single chain MHC class II molecules, each comprising alpha and
beta domain linked through amino acid linker and multimerization domain.
XX
PS Disclosure; Page 91-92; 147pp; English.
XX
CC The invention relates to a multimeric complex comprising a first
recombinant single chain major histocompatibility complex (MHC) class II
molecule and a second recombinant single chain MHC class II molecule,
each comprising an alpha domain and a beta domain linked through an
amino acid linker and a multimerization domain. The first and the second
molecule are linked through the multimerisation domain to form a
multimeric complex. The complex is useful for treating autoimmune
diseases. It is useful for treating insulin dependent diabetes, multiple
sclerosis, myasthenia gravis, pernicious anaemia, autoimmune
encephalomyelitis (EAE), rheumatoid arthritis and systemic lupus
erythematosus. The present sequence encodes a single chain MHC class II
molecule of the invention
XX
SQ Sequence 2346 BP; 560 A; 663 C; 646 G; 477 T; 0 U; 0 Other;
Query Match 48.9%; Score 726.2; DB 4; Length 2346;
Best Local Similarity 70.5%; Pred. No. 4.1e-185;
Matches 1095; Conservative 0; Mismatches 323; Indels 135; Gaps 4;
QY 68 GTGAAGACGACATTGAGCGGACCGAGTGGCTTCTATGTACAACTGTTTATCAGTCTC 127
|||||


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779 GTGAAGACGACATTGAGGCGGACCAAGTAGGCGTCTATGGTACAACTGTATATCAGTCTC 838
128 CTGGAGACATTGGCCAGATACACACATGAATTTGATGGTGATGAGTTGTTCTATGTGGACT 187
      |||
839 CTGGAGACATTGGCCAGTACACACATGAATTTGATGGTGATGAGTTGTTCTATGTGGACT 898
      |||
188 TGGATAAGAAGAAAACCTGCTGGAGGCTTCTGAGTTTGGCCAAATTGATACCTCTTTTGAGC 247
      |||
899 TGGATAAGAAGGAGACTACTGGATGCTTCTGAGTTTGGCCAAATTGACAAGCTTTTGACC 958
      |||
248 CCCAAGTTGGACTGCAAAACATAGCTGCAGAAAACACAACTTGGGAATCTTGACTAAGA 307
      |||
959 CCCAAGTTGGACTGCAAAACATAGCTGCAGAAAACATAGCTTGGAATCTTGACTAAGA 1018
      |||
308 GGTCAAAATTTCAACCCAGCTACCAATGAGGCTCTCAAGCGACTGTGTTCCCAAGTCCC 367
      |||
1019 GGTCAAAATTTCAACCCAGCTACCAATGAGGCTCTCAAGCGACTGTGTTCCCAAGTCCC 1078
      |||
368 GTGTGCTGTGGGTGAGCCCAACACCCTTATCTGCTTTGTGGAACAACATCTTCCACCTG 427
      |||
1079 GTGTGCTGTGGGTGAGCCCAACACCCCTCATCTGCTTTGTGGAACAACATCTTCCCTCCTG 1138
      |||
428 TGATCAACATCATGCTGCTCAGAAATAGCAAGTCAAGTCAAGCGCGCTTTATGAGACCA 487
      |||
1139 TGATCAACATCATGCTGCTCAGAAATAGCAAGTCAAGTCAAGCGCGCTTTATGAGACCA 1198
      |||
488 GCTTCTCTGCTCAACGCTGACCAATCTCTTCCAAAGCTGTCTTATCTCACTTTCATCCCTT 547
      |||
1199 GCTTCTCTGCTCAACGCTGACCAATCTCTTCCCAAGCTGTCTTATCTCACTTTCATCCCTT 1258
      |||
548 CTGATGATGACATTTATGATGCAAGGTGGAGCACTGGGGCTGTGAGAGCGCGTTCTGA 607
      |||
1259 CTGACGATGATATTTATGACTGCAAGGTGGAGCACTGGGGCTGTGAGAGCGCGTTCTGA 1318
      |||
608 AACACTGGG-----AA 618
      |||
1319 AACACTGGGCTAGCGAGGGGCGGGAAGCGGCGAGGGGAGCCAAACGACACCCCAT 1378
      |||
619 CTTGAGATTTCCAGCCCTCATGTGACAGCTGACAGAA----- 654
      |||
1379 CTGCTATCTACCTGCGCCCTGATCTGCTGCCAAACTTAATCTCATGTGTGACCTGGAT 1438
      |||
655 -ACTGAGGTGGAGATCCATACATGAGCTCCATGAGTCTGAGTCTGAGAAAGAGCTCCAGGC 713
      |||
1439 GCCTGCTCAAGGGCTATTTCCCTGAGCCAGTGACAGTCACTGGAATCTCTGGATCCCTGT 1498
      |||
714 CTTGGGAAGGAAT-----GCACAG 735
      |||
1499 CCAGCGGTGTGCACACTTCCAGCTGCTCTGAGTCTGACCTCTACACTGTGAGCAGCT 1558
      |||
736 CTGGAATGGGAGTTGCAAGCACTGGAAGAACTGGCTCAGGCAGCATCTGAGCCGAGA 795
      |||
1559 CAGTGACTGTCCCTCCAGCACTGGCCGAGCGAGACCGTCACTGCAACGTTGCCACC 1618
      |||
796 GGGCCACAAATCAAGCCCTGTCTCTCATGCAAAATGCCAG----- 835
      |||
1619 CGGCCAGCAGCAGCAAGGTGGAACAAGAAATTTGCGCCAGGATGTTGGTTGTAAGCCTT 1678
      |||
836 ---CACTTAACCTCTTGGGTGACCATCCGCTTCTATCTTCCCTCCAAAGATCAAGGATG 892
      |||
1679 GATATGTTACAGTCCAGGAATATCATCTGCTTTCATCTTCCCTCCAAAGGCCAAGGATG 1738
      |||
893 TACTCATGATCTCCCTGAGCCCATAGTCAATGTGTGTTGGTGGATGTGAGCGAGGATG 952
      |||
1739 TGCTCACCATTACTCTGACTCTCAAGTCAAGTGTGTTGTTGTGTAGACATCAGCAAGGATG 1798
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953 ACCCAGATGTCAGATCAGCTGGTTTGTGAACAAAGTGAAGTACACAGCTCAGACAC 1012
      |||
1799 ATCCGAGGTCCAGTTTCACTGGTTGTAGATGATGTGGAGGTGCACACAGCTCAGACGC 1858
      |||
1013 AAACCCATAGAGGATTAACAACAGTACTCTCCGGGTGCTCAGTGCCTCCCTCCCATCCAGC 1072
      |||
1859 AACCCCGGAGGAGCGATTTCAACAGCACTTTCGGCTCAGTCACTTCCCATCATGCT 1918
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```
QY 1073 ACCAGGACTGATGAGTGGCAAGGAGTTCAAATGCAAGGTCAACAAAGACCTCCACG 1132
      |||
Db 1919 ACCAGGACTGCTCAATGGCAAGGAGTTCAAATGCAAGGTCAACAGTGCAGCTTTCCCTG 1978
      |||
QY 1133 CGCCCATCGAGAACCATCTCAAAACCCAAAGGTCAGTAAGAGCTCCACAGGTATATG 1192
      |||
Db 1979 CCCCATCGAGAAACCATCTCCAAACCAAAGGCGAGHCCGAAGGCTCCACAGGTGTACA 2038
      |||
QY 1193 TCTTGCCTCCACCAAGAAGAGATGACTAAGAAAACAGGTCACTCTGACCTGCATGGTCA 1252
      |||
Db 2039 CCATTCCACCTCCCAAGGAGCAGATGGCCAAAGGATAAAGTCACTGACCTGCATGATAA 2098
      |||
QY 1253 CAGACTTCATCCCTGGAAGACATTTACGTGGAGTGGACCAACAAACGGGAAAACAGAGCTAA 1312
      |||
Db 2099 CAGACTTCTTCCCTGGAAGACATTTACGTGGAGTGGCAGTGGAAATGGGCGAGCCGCGAGA 2158
      |||
QY 1313 ACTACAGAACACTGMAACCACTCTGGACTCTGATGGTTCTTACTTCTATGATACAGCAAGC 1372
      |||
Db 2159 ACTCAAGAACACTCAGCCCATATGGACACAGATGGCTTACTTCTGTTTACATG 2218
      |||
QY 1373 TGAGAGTGGAAAAGAAAGAACTGGGTGGAAAAGAAATAGCTACTCTCTGTTTCACTGGTCCACG 1432
      |||
Db 2219 TCAATGTGCAGAGCAACTGGGAGGCGAGGAATACTTTCACTGCTCTGTGTTTACATG 2278
      |||
QY 1433 AGGCTCTGCACAATCAACACAGCACTAAGAGCTTCTCCCGGACTCCGGGTAAA 1485
      |||
Db 2279 AGGCGCTGCACACACCACCATACTGAGAAGAGCCTCTCCCACTCTCTCTGTTAAA 2331
      |||
```

RESULT 7

```
ABI99033
ID ABI99033 standard; cDNA; 2343 BP.
XX AC ABI99033;
XX
XX 25-FEB-2002 (first entry)
XX
XX MBP 90-101 CH1.H.CH2.CH3 coding sequence.
```

Mouse; MHC; major histocompatibility complex; MHC class II; multimer; single chain; immunosuppressive; antidiabetic; antiinflammatory; antianemic; antirheumatoid; antiarthritic; neuroprotective; vaccine; autoimmune disease; insulin dependent diabetes; multiple sclerosis; myasthenia gravis; pernicious anaemia; autoimmune encephalomyelitis; rheumatoid arthritis; systemic lupus erythematosus; ss.

Mus sp.
Synthetic.

WO200170245-A1.

27-SEP-2001.

22-MAR-2001; 2001WO-US009616.

22-MAR-2000; 2000US-0191274P.

15-MAY-2000; 2000US-020429P.

23-JAN-2001; 2001US-0264003P.

(CORI-) CORIXA CORP.

Carter D, Zhu S, Arimilli S, Wang A;

WPI; 2001-616371/71.

P-PSDB; ABB56463.

Multimeric complex for treating autoimmune diseases, comprises first and second single chain MHC class II molecules, each comprising alpha and beta domain linked through amino acid linker and multimerization domain.

Disclosure; Page 96; 147pp; English.

The invention relates to a multimeric complex comprising a first recombinant single chain major histocompatibility complex (MHC) class II molecule and a second recombinant single chain MHC class II molecule, each comprising an alpha domain and a beta domain linked through an amino acid linker and a multimerisation domain. The first and the second molecule are linked through the multimerisation domain to form a multimeric complex. The complex is useful for treating autoimmune diseases. It is useful for treating insulin dependent diabetes, multiple sclerosis, myasthenia gravis, pernicious anaemia, autoimmune encephalomyelitis (EAE), rheumatoid arthritis and systemic lupus erythematosus. The present sequence encodes a single chain MHC class II molecule of the invention

Sequence 2343 BP; 562 A; 665 C; 635 G; 481 T; 0 U; 0 Other;

Query Match 48.7%; Score 723.2; DB 4; Length 2343;

Best Local Similarity 70.4%; Pred. No. 2.6e-184;

Matches 1095; Conservative 0; Mismatches 323; Indels 138; Gaps 4;

```

68 GTGAAGACGACATTGAGGCGGACGACGATGGCTTCTATGCTACAACTGTTTATCAGTCTC 127
773 GTGAAGACGACATTGAGGCGGACGACGATGGCTTCTATGCTACAACTGTTTATCAGTCTC 832
128 CTGGAGACATTGGCCAGTACACACATGAATTTGATGGTGATGAGTTGTTCTATGTGGACT 187
833 CTGGAGACATTGGCCAGTACACACATGAATTTGATGGTGATGAGTTGTTCTATGTGGACT 892
188 TGGATAAGAAGAAACTGTCTGGAGGCTTCTGAGGCTTCTGAGTTTGGCCAAATGATACTCTTGGAGC 247
893 TGGATAAGAAGGAGACTATCTGGATGCTTCTGAGTTTGGCCAAATGATACTCTTGGAGC 952
248 CCCAAGGTGACCTGCAAAACATAGCTGCAGAAACACAACTTGGGAATCTTGACTAAGA 307
953 CCCAAGGTGACCTGCAAAACATAGCTGCAGAAACATAGCTGCAGAAACATCTTGACTAAGA 1012
308 GGTCAAAATTTCAACCCAGCTACCAATGAGGCTCCTCAAGCGACTGTGTTCCCAAGTCCC 367
1013 GGTCAAAATTTCAACCCAGCTACCAATGAGGCTCCTCAAGCGACTGTGTTCCCAAGTCCC 1072
368 GTGTGCTGTGGGTGAGCCCAACACCTTATCTGCTTTGTGTGACAAACATCTTCCCACTGG 427
1073 GTGTGCTGTGGGTGAGCCCAACACCTTATCTGCTTTGTGTGACAAACATCTTCCCTCTGG 1132
428 TGATCAACATCATCATGGCTCAGAAATAGCAATAGCAATGCAAGCGCGGTTTATGAGACCA 487
1133 TGATCAACATCATCATGGCTCAGAAATAGCAATAGCAATGCAAGCGCGGTTTATGAGACCA 1192
488 GCTTCTCTGTCAACCGTGACCAATTCCTTCCCAAGCTGTCTTATCTCACCTTTCATCCCTT 547
1193 GCTTCTCTGTCAACCGTGACCAATTCCTTCCCAAGCTGTCTTATCTCACCTTTCATCCCTT 1252
548 CTGATGATGACATTTATGATGCTGAAGGTGGAGACATGGGGGCTTGGAGGAGCGGTTCTGA 607
1253 CTGAGCATGATATTTATGATGCTGAAGGTGGAGACATGGGGGCTTGGAGGAGCGGTTCTGA 1312
608 AACACTGGG----- 616
1313 AACACTGGGCTAGCGAGGGGGGGAAGCGGGGAGAGAGTTAGCCAAACGACACGCC 1372
617 -AACCTGAGATTCCAGCCCTCATGTCTCAGAGCTGACAGAA----- 654
1373 CATCTGTCTATCCATCTGCGCCCTGGATCTGCTGCCCAACTCACTCCATGGTGACCTGG 1432
655 ----ACTGAGGTGGAGGATCCATACAGCTCCATCAGCTCAGCTGCGAAAGAGCTCA 710
1433 GATGCTGTGTCAGGGCTATTTCCCTGAGCCAGTGCAGTGCACCTGGAACCTCTGGATCCC 1492
711 GCGCTGGAGAGGAAAT-----GCA 732
1493 TGTCCAGCGGTGTGCACACCTTCCCAAGCTGTCTGACGCTTACACTCTTGAGCA 1552
733 CAGCTGGGAATGGAGTGTGCAAGCACTTGGAAAGGAAGAACTGGCTCAGGCGAGCATCTGAGGCC 792

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Db 1553 GCTCAGTGACTGTCCTCTCCAGCACTCGGCCCGAGGAGACCGTCACTGCAACGTTGCGCC 1612
Qy 793 AGAGGGCCCAATCAGCCCTGTCTCTCATGCAATGCCCCAG----- 835
Db 1613 ACCCGGCCAGCAGCAGCAAGGTGGAGCAAGAAATTTGTGCCCGAGGATTTGGTTGTAAGC 1672
Qy 836 -----CACCTAACTCTTTGGGTGGACCATCGCTCTTCATCTTCCCTCCAAAGATCAAGG 889
Db 1673 CTTGCATATGATACGATCCCGAGAGTATCATCTGTCTTCATCTTCCCGCCCAAGCCCAAGG 1732
Qy 890 ATGTACTCATGATCTCCCTGAGCCCATAGTCAATGTGTGTGGTGGATGTGAGCGAGG 949
Db 1733 ATGTGCTCACCATTACTCTGACTCTCAAGGTCAAGGTGTCAGTGTGTGTGGTAGACATCAGCAAGG 1792
Qy 950 ATGACCCAGATGTCAGATCAGCTGGTTTGTGAACAACTGGAAGTACACAGCTCAGA 1009
Db 1793 ATGATCCCGAGGTCCAGTTTCACTGGTTGTAGATGATGGAGGTGCACACAGCTCAGA 1852
Qy 1010 CACAAACCCATAGAGAGGATTACACAGTACTCTCCGGGTGGTCACTGCGCTCCCATCC 1069
Db 1853 CGCAACCCCGGAGAGGAGTTCACAGCACTTTCCGCTCAGTCACTGAGTGAATCTCCATCA 1912
Qy 1070 AGCACAGGACTGTGATGATGGCAAGGATTCAAATGCAAGGTCAACAAAGAGCTCC 1129
Db 1913 TGCACAGGACTGGCTCAATGGCAAGGATTCAAATGCAAGGTCAACAGTGCAGCTTCC 1972
Qy 1130 GAGCGCCATCGAGAGAACCATCTCAAAACCCAAAGGTCAGTAAAGCTCCACAGGTAT 1189
Db 1973 TGTGCCCCCATCGAGAAACCATCTCCAAACCCAAAGGTCAGTAAAGCTCCACAGGTGT 2032
Qy 1190 ATGTCTTGTGCTCCACAGAAAGAGATGACTAAGAAACAGGTCACTCTGACCTGCATGG 1249
Db 2033 ACACATTTCCACTCTCCAGGAGAGATGGCCAGGATTAAGTCACTGACCTGCATGA 2092
Qy 1250 TCACAGACTTCATGCTGAGACATTTACGTGAGTGGACCAACAAACCGGAAACACAGAGC 1309
Db 2093 TAAACAGACTTCTCCCTGAGACATTTACTGTGGAGTGGCAATGGGAGTGGAGCCAGCG 2152
Qy 1310 TAAACTACAGAACACTGAAACAGTCTGAGTCTGAGTGTCTTACTTTCATGTACAGCA 1369
Db 2153 AGAATCTACAGAAACACTCAGCCCATCTGACACAGATGGCTCTTACTTCTGCTACAGCA 2212
Qy 1370 AGCTCAGAGTGAAGAAAGAACTGGGTGGAAGAAATAGCTACTCTGTTCACTGCTGCTCC 1429
Db 2213 AGCTCAATGTGCAAGAGCAACTGGGAGGAGGAAATACTTTCACCTGCTGCTGTAC 2272
Qy 1430 ACGAGGCTTGCAATACCAACAGACTCAAGAGCTTCTCCCGGACTCCGGGTAAA 1485
Db 2273 ATGAGGCTTGCAACACCACCATCTGAGAGAGGCTCTCCCACTCTCCTGCTGATAA 2328

```

RESULT 8

AD007566

ID AD007566 standard; DNA; 1045 BP.

XX AC AD007566;

XX DT 15-JUL-2004 (first entry)

XX DE Fusion protein WT-IL-15-mIgG2a coding sequence.

XX KW immunosuppressive; antirheumatic; antiarthritic; antidiabetic;

XX KW neuroprotective; antipsoriatic; dermatological; antiinflammatory;

XX KW cytostatic; interleukin-15; immunoglobulin G; db; gene.

XX OS Synthetic.

XX OS Unidentified.

XX FH Key

XX CDS Location/Qualifiers

FT 1..1044

FT /*tag= a

FT /product= "WT-IL-15-mIgG2a"

FT /transl_except

/note= "(pos: 34. .36, aa: Thr"
/partial
/note= "no start codon"

WO2004035622-A2.

29-APR-2004.

13-OCT-2003; 2003WO-CH000666.

14-OCT-2002; 2002EP-00022869.

(HOFF) HOFFMANN LA ROCHE & CO AG F.

Dreher I, Moll T;

WPI; 2004-357203/33.
P-PSDB; ADO07559.

New fusion protein of interleukin-15 and Fc fragment, useful for treating e.g. transplantation disorders, autoimmune diseases and tumors, also related nucleic acid.

Disclosure; Fig 5; 63pp; German.

The present invention relates to a fusion protein consisting of wild-type interleukin-15 (IL-15) and an immunoglobulin G (IgG) Fc fragment, other than a murine IgG2b Fc fragment. The fusion proteins and coding sequences are used to prevent or treat consequences of transplantation and/or autoimmune diseases, e.g. rheumatoid arthritis, diabetes, multiple sclerosis, psoriasis, neurodermatitis, ulcerative colitis, tumours and AIDS, etc., and tissues or organs that express the protein are useful for transplantation into humans or other mammals, as allo-, auto- or xeno-transplants. Also transgenic animals that express the fusion proteins are useful as source of cells, tissues and organs for transplantation or to screen for pharmaceuticals and/or to identify toxic substances. The present sequence is a polypeptide coding sequence used in the exemplification of the invention.

Sequence 1045 BP; 331 A; 238 C; 235 G; 241 T; 0 U; 0 Other;

ery Match 47.2%; Score 701.2; DB 12; Length 1045;
st Local Similarity 93.3%; Pred. No. 1.6e-178;
tches 733; Conservative 0; Mismatches 53; Indels 0; Gaps 0;

700 AAAGAGCTCCAGCCCTGAGAGGAATGACAGCTGGATGGAGTTCGAACACTG 759
256 AAAGAATGTGAGGAATCGAGGAAAAAATATTAAAGAAATTTTGCAGAGTTTGTACAT 315
760 GAAAGGAATCGCTCAGGACGATCTGAGCCGAGAGGCCCAATCAAGCCCTGTCT 819
316 ATTGTCCAAATTTTCAACACTTGGATCCAGAGGGCCCAATCAAGCCCTGTCT 375
820 CCATGCAATGCCAGCAGCTTAACCTCTTGGGTGGACCAATCCGTCTTCACTTCCCTCA 879
376 CCATGCAATGCCAGCAGCTTAACCTCTTGGGTGGACCAATCCGTCTTCACTTCCCTCA 435
880 AAGATCAGGATGTACTCATGATCTCCCTGAGCCCAATGATGATGATGATGATGAT 939
436 AAGATCAGGATGTACTCATGATCTCCCTGAGCCCAATGATGATGATGATGATGAT 495
940 GTGAGGAGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 999
496 GTGAGGAGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 555
1000 ACAGCTCAGACACAAACCCATAGAGAGGATTACACAGTACTCTCCGGTGTCTAGTCCC 1059
556 ACAGCTCAGACACAAACCCATAGAGAGGATTACACAGTACTCTCCGGTGTCTAGTCCC 615
1060 CTCCTCCATCCAGACAGGATGATGATGATGATGATGATGATGATGATGATGATGAT 1119
616 CTCCTCCATCCAGACAGGATGATGATGATGATGATGATGATGATGATGATGATGAT 675

QY	1120	AAAGACCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTCAGTAAGAGCT	1179
Db	676	AAAGACCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTCAGTAAGAGCT	735
QY	1180	CCACAGGTATATGTTCTTGCCTCCACAGAGAGAGATGACTTAAGAAACAGTCACTCTG	1239
Db	736	CCACAGGTATATGTTCTTGCCTCCACAGAGAGAGATGACTTAAGAAACAGTCACTCTG	795
QY	1240	ACCTGCATGGTTCACAGACTTTCATGCTGGAACATTTACGTGGAGTGGACCAACCGG	1299
Db	796	ACCTGCATGGTTCACAGACTTTCATGCTGGAACATTTACGTGGAGTGGACCAACCGG	855
QY	1300	AAAAACAGAGCTAAACTACAGAACCACTGCTCTGGAACCTGATGATGATGATGATGAT	1359
Db	856	AAAAACAGAGCTAAACTACAGAACCACTGCTCTGGAACCTGATGATGATGATGATGAT	915
QY	1360	ATGTACAGCAGCTGAGAGTGGAAAGAGAACTGGGTGGAAAGAAATAGTACTCTCTGT	1419
Db	916	ATGTACAGCAGCTGAGAGTGGAAAGAGAACTGGGTGGAAAGAAATAGTACTCTCTGT	975
QY	1420	TCAGTGGTCCACGAGGGTCTGCACAATCACCACACGACTTAAGAGCTTCTCCGGACTCCG	1479
Db	976	TCAGTGGTCCACGAGGGTCTGCACAATCACCACACGACTTAAGAGCTTCTCCGGACTCCG	1035
QY	1480	GGTAAA 1485	
Db	1036	GGTAAA 1041	

RESULT 9
ADO07578
ID ADO07578 standard; DNA; 1108 BP.
XX ADO07578;
AC ADO07578;
XX 15-JUL-2004 (first entry)
XX Fusion protein coding sequence fragment 149-Fc.
XX immunosuppressive; antirheumatic; antiarthritic; antidiabetic;
KW neuroprotective; antipsoriatic; dermatological; antiinflammatory;
KW cystostatic; interleukin-15; immunoglobulin G; ds; gene; human.
OS Synthetic.
OS Unidentified.
XX WO2004035622-A2.
PN 29-APR-2004.
XX 13-OCT-2003; 2003WO-CH000666.
XX 14-OCT-2002; 2002EP-00022869.
XX (HOFF) HOFFMANN LA ROCHE & CO AG F.
XX Dreher I, Moll T;
XX WPI; 2004-357203/33.
XX New fusion protein of interleukin-15 and Fc fragment, useful for treating e.g. transplantation disorders, autoimmune diseases and tumors, also related nucleic acid.
XX Disclosure; Fig 11; 63pp; German.
XX The present invention relates to a fusion protein consisting of wild-type interleukin-15 (IL-15) and an immunoglobulin G (IgG) Fc fragment, other than a murine IgG2b Fc fragment. The fusion proteins and coding sequences are used to prevent or treat consequences of transplantation and/or autoimmune diseases, e.g. rheumatoid arthritis, diabetes, multiple sclerosis, psoriasis, neurodermatitis, ulcerative colitis, tumours and AIDS, etc., and tissues or organs that express the protein are useful for

transplantation into humans or other mammals, as allo-, auto- or xeno-transplants. Also transgenic animals that express the fusion proteins are useful as source of cells, tissues and organs for transplantation or to screen for pharmaceuticals and/or to identify toxic substances. The present sequence is a coding sequence used in the exemplification of the invention.

Sequence 1108 BP; 342 A; 255 C; 253 G; 257 T; 0 U; 0 Other;

Query Match 47.2%; Score 701.2; DB 12; Length 1108;

Best Local Similarity 93.3%; Pred. No. 1.7e-178;

Matches 733; Conservative 0; Mismatches 53; Indels 0; Gaps 0;

700 AAGAGCTCCAGGCCCTGGAGAGGAAATGACAGCTGGAATGGAGTTGCAAGCACTG 759

319 AAGAATGTGAGGAATGGAGGAAATAATTAAGAATTTTGGACATTTGTACAT 378

760 GAAAGGAATCTGGCTCAGGCAGCATCTGAGCCAGAGGGCCCAATCAAGCCCTGTCT 819

379 ATTGTCCAAATGTTTCATCAACACTTCGGATCCAGAGGGCCCAATCAAGCCCTGTCT 438

820 CATGCAAAATGCCAGCACTTAACCTCTTGGGTGGACCATCCGCTTTCATCTTCCCTCCA 879

439 CCATGCAAAATGCCAGCACTTAACCTCTTGGGTGGACCATCCGCTTTCATCTTCCCTCCA 498

880 AAGATCAAGGATCTACTCATGATCTCCCTGAGCCCATAGTCACATGTGTGGTGGAT 939

499 AAGATCAAGGATCTACTCATGATCTCCCTGAGCCCATAGTCACATGTGTGGTGGAT 558

940 GTGAGGAGGATGACCCAGATGTCAGATCAGCTGGTTGTGAAACAGTGGGAAGTACAC 999

559 GTGAGGAGGATGACCCAGATGTCAGATCAGCTGGTTGTGAAACAGTGGGAAGTACAC 618

1000 ACAGCTCAGACACAAACCCATAGAGAGGATTACAAGTACTCTCCGGGTGGTCAAGTCC 1059

619 ACAGCTCAGACACAAACCCATAGAGAGGATTACAAGTACTCTCCGGGTGGTCAAGTCC 678

1060 CTCCCATCCAGCAGGACTGGATGAGTGGCAAGGATTTCAATGCAAGTCAACAC 1119

679 CTCCCATCCAGCAGGACTGGATGAGTGGCAAGGATTTCAATGCAAGTCAACAC 738

1120 AAGACTCTCCAGGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTCAGTAAGAGCT 1179

739 AAGACTCTCCAGGCCCATCGAGAGAACCATCTCAAAACCCAAAGGGTCAGTAAGAGCT 798

1180 CCAAGGATATCTTTGCTTCCACCAAGAAAGAGATGATTAAGAAACAGGTCATCTTG 1239

799 CCAAGGATATCTTTGCTTCCACCAAGAAAGAGATGATTAAGAAACAGGTCATCTTG 858

1240 ACCTGCATGCTCAGACTTCATGCTGAGACATTTAGTGGAGTGGACCAACACGGG 1299

859 ACCTGCATGCTCAGACTTCATGCTGAGACATTTAGTGGAGTGGACCAACACGGG 918

1300 AAAAAAGAGCTAACTACAAGAACACTGAACCACTCTGACTCTGATGGTTCTTACTTC 1359

919 AAAAAAGAGCTAACTACAAGAACACTGAACCACTCTGACTCTGATGGTTCTTACTTC 978

1360 ATGTAAGAGCTGAGAGTGGAAAAAGAACTGGGTGGAAGAAATAGTACTCTCTGT 1419

979 ATGTAAGAGCTGAGAGTGGAAAAAGAACTGGGTGGAAGAAATAGTACTCTCTGT 1038

1420 TCAGTGGTCCAGAGGCTGTCACATCACCACAGCTTAAGAGCTTCTCCGGAATCCG 1479

1039 TCAGTGGTCCAGAGGCTGTCACATCACCACAGCTTAAGAGCTTCTCCGGAATCCG 1098

1480 GGTAAA 1485

1099 GGTAAA 1104

ULT 10

:07577

ADO07577 standard; DNA; 1108 BP.

XX

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1000 ACAGCTCAGACACAAACCATAGAGGATTACACAGTACTCTCCGGTGGTCAGTGCC 1059
619 ACAGCTCAGACACAAACCATAGAGGATTACACAGTACTCTCCGGTGGTCAGTGCC 678
1060 CTCCCATCCAGCACACAGGACTGGATGAGTGCCAAAGGATTCAAAATGCAAGGTCAACAC 1119
679 CTCCCATCCAGCACACAGGACTGGATGAGTGCCAAAGGATTCAAAATGCAAGGTCAACAC 738
1120 AAAGACTCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGTCAGTAAGCT 1179
739 AAAGACTCTCCAGCGCCCATCGAGAGAACCATCTCAAAACCCAAAGGTCAGTAAGCT 798
1180 CCACAGGTATATGTCTTGCTCCACAGAGAGAGATGACTAAGAAACAGGTCACTCTG 1239
799 CCACAGGTATATGTCTTGCTCCACAGAGAGAGATGACTAAGAAACAGGTCACTCTG 858
1240 ACCTGCATGGTCCACAGACTTCATGCTCTGAAGACATTTACGTGGAGTGCACCAACACGG 1299
859 ACCTGCATGGTCCACAGACTTCATGCTCTGAAGACATTTACGTGGAGTGCACCAACACGG 918
1300 AAAACAGAGCTAACTACAGAAACACTGAACAGTCTCGGACTCTGATGGTCTTACTTC 1359
919 AAAACAGAGCTAACTACAGAAACACTGAACAGTCTCGGACTCTGATGGTCTTACTTC 978
1360 ATGTACAGCAAGCTGAGAGTGGAAGAAAGAACTGGTGGAAGAAATAGCTACTCTGT 1419
979 ATGTACAGCAAGCTGAGAGTGGAAGAAAGAACTGGTGGAAGAAATAGCTACTCTGT 1038
1420 TCAGTGGTCCACAGGGTCTGCACAAATCAACACAGACTAAGAGCTTCTCCGGACTCGG 1479
1039 TCAGTGGTCCACAGGGTCTGCACAAATCAACACAGACTAAGAGCTTCTCCGGACTCGG 1098
1480 GSTAAA 1485
1099 GGTAAG 1104

LT 11

5694

ADL15694 standard; DNA; 990 BP.

ADL15694;

20-MAY-2004 (first entry)

Murine immunoglobulin heavy chain constant region DNA SeqID 68.

mouse; murine; antibody; gene; ds; beta-amyloid; A-beta;
amyloid beta A4 precursor protein; APP; presenilin;
lipoprotein receptor related protein; LRP; beta-amyloid 42; A-beta 42;
Alzheimer's disease; neuroprotective; nontropic.

Mus musculus.

WO2004018997-A2.

04-MAR-2004.

20-AUG-2003; 2003WO-US026173.

20-AUG-2002; 2002US-0405417P.

18-SEP-2002; 2002US-0411974P.

(NEUR-) NEUROGENETICS INC.

Kounnas M, Patrick A, Velicelebi G, Wagner S;

WPI; 2004-226902/21.

P-PSDB; ADL15695.

New polypeptide comprises a sequence of amino acids that is selectively
reactive with beta-amyloid peptide 42 or at least one complementarity-

PT determining region of antibody A387 or B436, useful for treating
PT Alzheimer's disease.
XX
PS Disclosure; SEQ ID NO 68; 408pp; English.
XX
CC This invention relates to novel methods and compositions for detecting
CC and modulating beta-amyloid (A-beta) peptide levels and the processing of
CC amyloid beta A4 precursor protein (APP). Specifically, it refers to
CC methods of assessing the presenilin activity of compounds using the
CC lipoprotein receptor related protein (LRP), in order to identify
CC presenilin proteins that can be used to affect the processing of APP. The
CC present invention describes methods to identify agents that modulate
CC presenilin activity and A-beta levels, in particular beta-amyloid 42 (A-
CC beta 42), such that the agent is selectively reactive with A-beta 42 and
CC binds at least one complementarity determining region (CDR) of either
CC antibody A387 or antibody B436. As such, the polypeptides, nucleic acids
CC and antibodies are useful for treating Alzheimer's disease, accordingly
CC the compositions exhibit neuroprotective and nontropic activities. This
CC polynucleotide sequence is a murine antibody chain DNA fragment of the
CC invention.
XX
SQ Sequence 990 BP; 274 A; 286 C; 235 G; 195 T; 0 U; 0 Other;
Query Match 47.2%; Score 700.6; DB 12; Length 990;
Best Local Similarity 100.0%; Pred. No. 2.3e-178;
Matches 700; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 786 TGAGCCAGAGGGCCCAATCAAGCCCTGCTCCATGCAAAATCCCGACACCTAACT 845
Db 291 TGAGCCAGAGGGCCCAATCAAGCCCTGCTCCATGCAAAATCCCGACACCTAACT 350
QY 846 CTGGGTGGACCATCCGTTTCATCTTCCCTCCAAAGATCAAGGATGACTCATGATCTC 905
Db 351 CTGGGTGGACCATCCGTTTCATCTTCCCTCCAAAGATCAAGGATGACTCATGATCTC 410
QY 906 CTGAGCCCCATAGTCACATGTGTGGTGGATGAGCGAGGATGACCCAGATGTCCA 965
Db 411 CTGAGCCCCATAGTCACATGTGTGGTGGATGAGCGAGGATGACCCAGATGTCCA 470
QY 966 GATCAGCTGGTTTGTGAACAAACGTGGAAGTACACACAGCTCAGACACAAACCCATAGAGA 1025
Db 471 GATCAGCTGGTTTGTGAACAAACGTGGAAGTACACACAGCTCAGACACAAACCCATAGAGA 530
QY 1026 GGATTACAAAGTACTCTCCGGGTGGTCAAGTGCCTCCCTCCATCCAGCACCAGGACTGGAT 1085
Db 531 GGATTACAAAGTACTCTCCGGGTGGTCAAGTGCCTCCCTCCATCCAGCACCAGGACTGGAT 590
QY 1086 GAGTGGCAAGGATTCAAATGCNAGGTCAACAAACAAAGACCTCCCGAGCCCATCGAGAG 1145
Db 591 GAGTGGCAAGGATTCAAATGCNAGGTCAACAAACAAAGACCTCCCGAGCCCATCGAGAG 650
QY 1146 AACCATCTCAAAACCCAAAGGTCAGTAAGAGCTCCACAGGTATATGCTTCCCTCCACC 1205
Db 651 AACCATCTCAAAACCCAAAGGTCAGTAAGAGCTCCACAGGTATATGCTTCCCTCCACC 710
QY 1206 AGAAGAGAGATGACTAAGAAACAGGTCACTCTGACCTGCTGATGGTTCACAGACTCATGCC 1265
Db 711 AGAAGAGAGATGACTAAGAAACAGGTCACTCTGACCTGCTGATGGTTCACAGACTCATGCC 770
QY 1266 TGAAGACATTTACGTGGAGTGCACCAACGCGGAAACACAGGCTAAACTACAAGAACAC 1325
Db 771 TGAAGACATTTACGTGGAGTGCACCAACGCGGAAACACAGGCTAAACTACAAGAACAC 830
QY 1326 TGAACCACTCTGGACTCTGATGGTTCCTTACTTCACTAGTACAGAGCTGAGAGTGA 1385
Db 831 TGAACCACTCTGGACTCTGATGGTTCCTTACTTCACTAGTACAGAGCTGAGAGTGA 890
QY 1386 GAAGAACTGGGTGGAAAGAAATAGTACTCTGTTTCACTGGTCCAGAGGGTCTGCACAA 1445
Db 891 GAAGAACTGGGTGGAAAGAAATAGTACTCTGTTTCACTGGTCCAGAGGGTCTGCACAA 950
QY 1446 TCACACACAGCTAGAGCTTCTCCCGGACTCCCGGTAAA 1485

951 TCACCACACGACTAAGAGCTTCTCCGGACTCCGGGTAA 990

:ULT 12
 :20762

AEC20762 standard: CDNA: 1401 BP.

AEC20762;

20-OCT-2005 (first entry)

M-CSF specific murine antibody RX1 heavy chain cDNA.

endocrine-gen.; antiarthritic; antibacterial; antiinflammatory; antihumetic; antithyroid; bone metastases; calcium antagonist; cancer; cardiovascular-gen.; degeneration; eating-disorders-gen.; endocrine disease; endocrine-gen.; endocrine-gen.; gastrointestinal-gen.; gene; genetic disorder; heavy chain; hepatotropic; hypercalcemia; immune disorder; immunotherapy; inflammation; monoclonal antibody; mouth disease; musculoskeletal disease; neoplasm; nephrotropic; osteopathic; osteopetrosis; osteoporosis; pages disease; periodontal disease; pharmacological; rheumatoid arthritis; ss.

Mus musculus.

WO2005068503-A2.

28-JUL-2005.

06-JAN-2005; 2005WO-US000546.

07-JAN-2004: 2004US-0535181P.

02-JUN-2004; 2004US-0576417P.

(CHIR) CHIRON CORP.

(XOMA) XOMA TECHNOLOGY LTD.

Liu C, Zimmerman DL, Harrowe GM, Koths K, Kavanaugh WM, Long L; Calderon-Cacia M, Horwitz AH;

WPI: 2005-597707/61.

P-PSDB: ABC20763.

Novel non-murine antibody that competes with monoclonal antibody RX1 for binding to macrophage colony stimulating factor, useful for treating hypogonadism, hypercalcemia, rickets, scurvy, homocystinuria, cancer, osteoporosis.

Claim 67: SEQ ID NO 1: 269pp: English.

The invention describes a non-murine antibody (I) that competes with monoclonal antibody RX1 for binding to macrophage colony stimulating factor (M-CSF) by more than 75%, where the monoclonal antibody RX1 has the heavy chain and light chain amino acid sequences having a fully defined 447 amino acids (SEQ ID No. 2) and 214 amino acids (SEQ ID No. 4) sequences given in the specification, respectively. (I) is useful for preventing a subject afflicted with a disease that causes or contributes to osteolysis, where the antibody effectively reduces the severity of bone loss associated with the disease. The disease is chosen from metabolic bone diseases associated with relatively increased osteoclast activity, including endorinopathies, hypercalcemia, deficiency states, chronic diseases, and hereditary diseases, cancer, osteoporosis, osteopetrosis, inflammation of bone associated with arthritis and rheumatoid arthritis, periodontal disease, fibrous dysplasia, and/or Paget's disease. (I) is useful for preventing or treating metastatic cancer. Antibodies of the invention are useful for preventing or reducing bone loss; osteolysis; metastatic cancer to bone and cancer. (I) is useful for manufacturing a medicament for preventing or reducing bone loss in a patient exhibiting osteolysis, manufacturing a medicament for treating a patient afflicted with a disease that causes or contributes to osteolysis, and metastatic cancer to bone in a patient suffering from metastatic cancer, for manufacturing a medicament for treating a patient having cancer. (II) in synergistic combination, is useful for preparing a

WO2005094846-A1.

13-OCT-2005.

30-MAR-2005; 2005WO-JP006189.

30-MAR-2004; 2004JP-00100649.

(RENO-) RENOMEDIX INST INC.

Fujinaga K, Shinagawa M, Niitsu Y, Hamada H, Horiuchi M;
Homnou O, Umetani A;

WPI; 2005-725409/74.

Agent useful for treating prion disease or delivering a substance to a
lesioned region of prion disease, comprises a mesenchymal cell.

Claim 4; SEQ ID NO 5; 34pp; Japanese.

The invention describes an agent (I) for treating prion disease or
delivering a substance to the lesioned region of prion disease,
comprising a mesenchymal cell. Also described are: a nucleic acid (II)
having an anti-prion antibody gene comprising: an antibody heavy chain
gene having SEQ ID No: 1, 3, 5, 30, 32 and 34, a nucleotide sequence
consisting of a degenerate genetic code, which encodes a polypeptide same
as that of the above nucleotide sequence, a nucleotide sequence, which is
a mutant of the above sequences, or a nucleotide sequence that is
complementary to the above sequences and that hybridizes under stringent
conditions with the above sequences; and an antibody light chain gene
having SEQ ID No: 2, 4, 6, 31, 33 and 35, a nucleotide sequence
consisting of degenerate genetic code, which encodes a polypeptide same
as that of the above nucleotide sequence, a nucleotide sequence, which is
a mutant of the above sequences, or a nucleotide sequence that is
complementary to the above sequences and that hybridizes under stringent
conditions with the above sequences; a vector (III) comprising (II); an
anti-prion chimeric antibody (IV) comprising variable region of antibody
encoded by (II) and constant region of antibody of animal other than
mouse; a nucleic acid that encodes (IV); preparing (M1) a cell having
abnormal prion proliferation inhibition activity, comprising transducing
a gene that provides abnormal prion proliferation inhibition activity to
the cell; a cell (V) having abnormal prion proliferation inhibition
activity, being obtainable by (M1) or by utilizing (II) or (III); a
sustainable formulation (VI) for discharge of an anti-prion antibody
utilized for treating prion disease; use of a mesenchymal cell for
producing an agent for delivering a substance to the lesioned region of
prion disease; and delivering a substance to the lesioned region of a
prion disease, comprising utilizing mesenchymal cell. (I) is useful for
treating prion disease or delivering a substance to the lesioned region
of prion disease. (II), (III) Or (M1) is useful for preparing a cell
having abnormal prion proliferation inhibition activity. (II), (III),
(III), (IV) Or (VI) is useful for treating prion disease. (I) Enables
improvement of the symptoms of prion disease. This sequence an anti-PrP
monoclonal antibody heavy chain. Note: This sequence does not appear in
the printed specification but has been obtained in electronic format
directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

Sequence 1560 BP; 413 A; 424 C; 388 G; 335 T; 0 U; 0 Other;

ery Match 47.2%; Score 700.6; DB 14; Length 1560;
st Local Similarity 100.0%; Pred. No. 2.8e-178;
tches 700; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

786 TGAGCCGAGGCGCCACAAATCAAGCCCTGCTCCATGCAATGCCAGCACCTAACT 845

850 TGAGCCGAGGCGCCACAAATCAAGCCCTGCTCCATGCAATGCCAGCACCTAACT 909

846 CTTGGGTGGACCATCGCTTTCATCTTCCCTCCAAAGATCAAGGATGACTCATGATCTC 905

910 CTTGGGTGGACCATCGCTTTCATCTTCCCTCCAAAGATCAAGGATGACTCATGATCTC 969

906 CCTGAGCCCCCATAGTCACATGTTGGTGGATGTGAGCGGAGGATGACCCAGATGTCCA 965

Db CCTGAGCCCCCATAGTCACATGTTGGTGGATGTGAGCGGAGGATGACCCAGATGTCCA 1029
QY GATCAGCTGGTTTGTGAACAACGTTGGAAGTACACACAGCTCAGACACAAACCCATAGAGA 1025
Db GATCAGCTGGTTTGTGAACAACGTTGGAAGTACACACAGCTCAGACACAAACCCATAGAGA 1089
QY GGATTACAAACAGTACTCTCTCCGGGGTGGTCAGTGCCTCCCATCCAGCACCAGGACTGGAT 1085
Db GGATTACAAACAGTACTCTCTCCGGGGTGGTCAGTGCCTCCCATCCAGCACCAGGACTGGAT 1149
QY GAGTGGCAAGGAGTTCAAAATGCAAGGTCACAAACAAGACCTCCAGCGGCCCATCGAGAG 1145
Db GAGTGGCAAGGAGTTCAAAATGCAAGGTCACAAACAAGACCTCCAGCGGCCCATCGAGAG 1209
QY AACCATCTCAAAACCCAAAGGGTCAGTAAAGAGTCCACAGGTATATGTCTTGCCTCCACC 1205
Db AACCATCTCAAAACCCAAAGGGTCAGTAAAGAGTCCACAGGTATATGTCTTGCCTCCACC 1269
QY AGAAGAAGAGATGACTTAAGAAACAGGTCATCTGACCTGTCATGGTCAACAGACTTCATGCC 1265
Db AGAAGAAGAGATGACTTAAGAAACAGGTCATCTGACCTGTCATGGTCAACAGACTTCATGCC 1329
QY TGAAGACATTTTACGTGGAGTGGACCAACACCGGNAACAGAGCTAAACTACAAGAACAC 1325
Db TGAAGACATTTTACGTGGAGTGGACCAACACCGGNAACAGAGCTAAACTACAAGAACAC 1389
QY TGAAGACATTTTACGTGGAGTGGACCAACACCGGNAACAGAGCTAAACTACAAGAACAC 1385
Db TGAAGACATTTTACGTGGAGTGGACCAACACCGGNAACAGAGCTAAACTACAAGAACAC 1449
QY TGAAGACATTTTACGTGGAGTGGACCAACACCGGNAACAGAGCTAAACTACAAGAACAC 1445
Db TGAAGACATTTTACGTGGAGTGGACCAACACCGGNAACAGAGCTAAACTACAAGAACAC 1509
QY TCACCACACGACTAAGAGCTTCTCCCGGACTCCCGGGTAAA 1485
Db TCACCACACGACTAAGAGCTTCTCCCGGACTCCCGGGTAAA 1549

RESULT 14

ADV26108 standard; DNA; 1569 BP.

AC ADV26108;

DT 10-MAR-2005 (first entry)

XX Mouse OKT3 VH gene.

DE ds; gene; immunostimulant; immunogenicity; antibody.

XX Mus sp.

XX WO2004108158-A1.

XX 16-DEC-2004.

XX 28-MAY-2004; 2004WO-US017219.

XX 02-JUN-2003; 2003US-0475155P.

XX (ALEX-) ALEXION PHARM INC.

XX Rother RP, Faas-Knight S, Wu D, Carr FJ, Hamilton A;

XX WPI; 2005-031597/03.

XX P-PSDB; ADV26107.

XX New de-immunized anti-CD3 antibody, useful for stimulating an immune
response against infections and for treating infections.
XX Disclosure; Fig 1a; 75pp; English.

924 GATCAGCTGGTTTGTGAACAACGTTGAAAGTACACACAGCTCAGACACAAACCCATAGAGA 983
1026 GGATTACAACAGTACTCTCCGGTGGTCAGTGCCTCCCATCCAGACACAGGACTGGAT 1085
984 GGATTACAACAGTACTCTCCGGTGGTCAGTGCCTCCCATCCAGACACAGGACTGGAT 1043
1086 GAGTGGCAAGGAGTTCAAATGCAAGGTCAACAACAAGACCTCCAGCGCCCATCGAGAG 1145
1044 GAGTGGCAAGGAGTTCAAATGCAAGGTCAACAACAAGACCTCCAGCGCCCATCGAGAG 1103
1146 AACCATCTCAAAAACCAAGGTCAGTAAGAGCTCCACAGGTATATGTCTTGCCTCCACC 1205
1104 AACCATCTCAAAAACCAAGGTCAGTAAGAGCTCCACAGGTATATGTCTTGCCTCCACC 1163
1206 AGAAGAAGAGATGACTAAGAACAGGTCACTCTGACCTGCATGGTCCACAGACTTCATGCC 1265
1164 AGAAGAAGAGATGACTAAGAACAGGTCACTCTGACCTGCATGGTCCACAGACTTCATGCC 1223
1266 TGAAGACATTTACGTGGAGTGGACCACAAACCGGAAACACAGAGCTAAACTACAAGAACAC 1325
1224 TGAAGACATTTACGTGGAGTGGACCACAAACCGGAAACACAGAGCTAAACTACAAGAACAC 1283
1326 TGAACCACTCTGGACTCTGATGGTTCTTAATTATGTACAGCAAGCTGAGAGTGGAAAA 1385
1284 TGAACCACTCTGGACTCTGATGGTTCTTAATTATGTACAGCAAGCTGAGAGTGGAAAA 1343
1386 GAAGAACTGGGTGGAAAGAAATAGCTACTCTGTTTCAGTGGTCCACAGAGGTCTGCACAA 1445
1344 GAAGAACTGGGTGGAAAGAAATAGCTACTCTGTTTCAGTGGTCCACAGAGGTCTGCACAA 1403
1446 TCACCACACGACTAAGAGCTTCTCCCGGACTCCGGGTAAA 1485
1404 TCACCACACGACTAAGAGCTTCTCCCGGACTCCGGGTAAA 1443

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Maximum Match 100%

Listing first 45 summaries

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2: Geneseqn1990s:*

3: Geneseqn2000s:*

4: Geneseqn2001as:*

5: Geneseqn2001bs:*

6: Geneseqn2002as:*

7: Geneseqn2002bs:*

8: Geneseqn2003as:*

9: Geneseqn2003bs:*

10: Geneseqn2003cs:*

11: Geneseqn2003ds:*

12: Geneseqn2004as:*

13: Geneseqn2004bs:*

14: Geneseqn2005s:*

15: Geneseqn2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Alt	Query	Score	Match	Length	DB	ID	Description
1	921	96.8	921	5	AAF55099		Aaf55099 DNA encod
2	709.2	74.6	893	2	AAT04262	PF	Aat04262 Hybrid IA
3	665.2	69.9	945	12	ADQ311225		Adq311225 I-Ab(beta
4	648	68.1	1013	2	AAT04269	PR	Aat04269 Hybrid IA
5	628.6	66.1	915	12	ADQ311228		Adq311228 I-Ab(beta
6	607.8	63.9	1382	2	AAT86989		Aat86989 SCEL sing
7	607.8	63.9	1382	8	ACA60744		Acac60744 Mouse MHC
8	607.8	63.9	1385	2	AAT86987	SSC1	Aat86987 SSC1 sing
9	607.8	63.9	1385	8	ACA60742		Acac60742 Mouse MHC
10	607.8	63.9	1508	2	AAT86988	SCT1	Aat86988 SCT1 sing
11	607.8	63.9	1508	2	AAT89069		Aat89069 Single ch
12	607.8	63.9	1508	8	ACA60743		Acac60743 Mouse MHC
13	606.2	63.7	1382	2	AAT17588		Aat17588 Vector SC
14	606.2	63.7	1385	2	AAT17586		Aat17586 Vector SS
15	606.2	63.7	1508	2	AAT17587		Aat17587 Vector SC
16	598.6	62.9	4724	2	AAV12068		Aav12068 Murine IA
17	561.8	59.1	798	12	ADJ75986		Adj75986 Marker ge
18	561.8	59.1	798	14	ADX26090		Adx26090 Novel cel

ALIGNMENTS

RESULT 1

AAF55099

ID AAF55099 standard; DNA; 921 BP.

AC AAF55099;

XX 15-MAY-2001 (first entry)

XX DNA encoding a fusion protein comprising a beta chain of MHC.

XX Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;

XX major histocompatibility complex; Fc region; antigen; T lymphocyte;

XX immunostimulant; vaccine; infection; tumour; ss.

XX Synthetic.

XX Key

XX Location/Qualifiers

FT CDS

FT i. 921

FT /*tag= a

XX WO200109194-A1.

XX 08-FEB-2001.

XX 28-JUL-2000; 2000WO-FR002193.

XX 29-JUL-1999; 99FR-00009862.

XX (CNRS) CNRS CENT NAT RECH SCI.

XX Glaichenhaus N, Malherbe L;

XX WPI; 2001-182944/18.

XX P-PSDB; AAB67481.

XX New soluble recombinant protein, useful e.g. as immunostimulant, comprises dimeric major histocompatibility complex molecule fused to immunoglobulin Fc region.

XX Example 1; Page 34-35; 43pp; French.

08-SEP-1995.
03-MAR-1995; 95WO-US002689.
04-MAR-1994; 94US-00207481.
(NAJE-) NAT JEWISH CENT IMMUNOLOGY & RESPIRATORY.
Kappler JW, Marrack P;
WPI; 1995-320543/41.
P-PSDB; AAR82533.
Peptide-MHC complex comprising antigenic peptide, linker and MHC segment
- useful as reagents for the treatment of diseases including auto-immune
diseases, immuno-stimulatory diseases or graft-host rejection.
Example 1; Page 53; 94pp; English.
This sequence represents a hybrid IA beta chain gene, containing the
chicken ovalbumin peptide (cOVA). This sequence was used in the
construction of a hybrid IA alpha beta dimer. The encoded protein (pIad-
OVA) was found to be more stable than the IA alpha beta dimer. The
stability was decreased by the addition of a MHC groove specific binding
peptide (e.g. see AAR82527, AAR82528 and AAR82531), compared to an
increase seen on the addition of a MHC binding peptide to IE k/d-MCC.
These complexes may be used to regulate an immune response. The complexes
are capable of being recognised by a TCR alone or in combination with
additional MHC proteins. These complexes are useful for therapeutic
purposes and experimental purposes. They can also be used as reagents for
the treatment of diseases including autoimmune diseases, immunodeficiency
diseases, immunoproliferation diseases, and graft-host rejection
Sequence 893 BP; 204 A; 239 C; 275 G; 175 T; 0 U; 0 Other;
Very Match 74.6%; Score 709.2; DB 2; Length 893;
%st Local Similarity 94.7%; Pred. No. 2.1e-155;
atches 761; Conservative 0; Mismatches 28; Indels 15; Gaps 2;
8 GGAATTTCTTAGAGATGGCTCTGCAGATCCCGAGCTCTCTCTCTCAGCTGCTGTGTGT 67
48 GGAATTTCTTAGAGATGGCTCTGCAGATCCCGAGCTCTCTCTCTCAGCTGCTGTGTGT 107
68 GCTGATGCTCTGAGCAGCCCGGGACTGAGGCGGGAATCTCATCTGCTTCGCGCTC 127
108 GCTGATGCTGCTGAGCAGCCCGGGACTGAGGCGGGAATCTCC-----GTACATGCTGCC 162
128 GCTGAGGACCCGATCGTGTGTCGGCAGCTGGGACGGAGTGGGGCTCACTAGTGCC 187
163 CATGCTGAGATCAATGAGGCTGGCAG-----AGGAGTGGGGGCTCACTAGTGCC 212
188 CCGAGGCTCTGAGAGTGGAGGCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTG 247
213 CCGAGGCTCTGAGAGTGGAGGCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTG 272
248 CTACTACACCAACGGGACGGGACATACGGCTCGTGACACAGATACATCTACACCGGA 307
273 CTACTACACCAACGGGACGGGACATACGGCTCGTGACACAGATACATCTACACCGGA 332
308 GGAGTACGTCGCTACGACGAGCGTGGGCGGAGTACCGCGGTGACCGAGTGGGGCG 367
333 GGAGTACGTCGCTACGACGAGCGTGGGCGGAGTACCGCGGTGACCGAGTGGGGCG 392
368 GCCAGACGCGGAGTACTGGAAACAGCCAGCGGAGATCTCTGAGCGGAACCGGCGCGAGGT 427
393 GCCAGACGCGGAGTACTGGAAACAGCCAGCGGAGATCTCTGAGCGGAACCGGCGCGAGGT 452
428 GGACACGGGCTGACACACAACTACGAGGGGCGGAGACGACGACCTCTCTGCGGGGCT 487
453 GGACACGGGCTGACACACAACTACGAGGGGCGGAGACGACGACCTCTCTGCGGGGCT 512
488 TGAACAGCCCAATGTGCGCCATCTCTCTGTCAGGACAGAGGCGCTCAACACCACACAC 547

Db 513 TGAAAGCCCAATGTGCGCCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCACACAC 572
QY 548 TCTGCTCTGTTCCGGTGAAGATTTCTAACCAGCCAGATCAAGTCCGCTGGTTTCAGGAA 607
Db 573 TCTGCTCTGTTCCGGTGAAGATTTCTAACCAGCCAGATCAAGTCCGCTGGTTTCAGGAA 632
QY 608 TGGCCAGGAGGAGACAGTGGGGGTCTCATCCACACAGCTTTATTAGGAATGGGACTGGAC 667
Db 633 TGGCCAGGAGGAGACAGTGGGGGTCTCATCCACACAGCTTTATTAGGAATGGGACTGGAC 692
QY 668 CTTCCAGGCTCTGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 727
Db 693 CTTCCAGGCTCTGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 752
QY 728 TGTGAGGATCCAGCTGAGAGCCCATCATCTGTGAGTGGAGGGGACAGTCCGAGTC 787
Db 753 TGTGAGGATCCAGCTGAGAGCCCATCATCTGTGAGTGGAGGGGACAGTCCGAGTC 812
QY 788 TGCCCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 811
Db 813 TGCCCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 836
RESULT 3
ADQ31225
ID ADQ31225 standard; cDNA; 945 BP.
XX
AC ADQ31225;
XX
DT 07-OCT-2004 (first entry)
XX
DE I-Ab(beta)-Cholera toxin B subunit-leucine zipper (LZ)-Bira fusion cDNA.
XX
KW class II major histocompatibility complex; MHC; CD4+ T-cell detection;
flow cytometry; mucous membrane invasive antigen;
KW I-Ab(beta)-Cholera toxin B subunit-leucine zipper-Bira fusion; CTB; ss;
KW gene.
XX
OS Vibrio cholerae.
Unidentified.
XX
FH Key Location/Qualifiers
FT CDS 1..945
FT /tag= a
FT /product= "I-Ab(beta)-Cholera toxin B subunit (CTB) -
leucine zipper (LZ)-Bira fusion cDNA"
XX
PN JP2004196789-A.
XX
PD 15-JUL-2004.
XX
PF 03-DEC-2003; 2003JP-00404367.
XX
PR 03-DEC-2002; 2002JP-00351818.
XX
PA (SENT-) SENTAN KAGAKU GIJUTSU INCUBATION CENT KK.
XX
DR WPI; 2004-546819/53.
DR P-PSDB; ADQ31224.
XX
PT Peptide-Class II major histocompatibility complex (MHC) composite, useful
for detecting antigen specific CD4+ T-cell, comprises antigen peptide
containing epitope of mucous membrane invasive protein, and extracellular
region of MHC.
XX
PS Example 1; SEQ ID NO 10; 30pp; Japanese.
XX
CC The invention relates to a novel class II major histocompatibility
complex (MHC) antigenic peptide composite comprising a peptide containing
the T-cell antigenic determinant of a mucous membrane invasive protein
and the extracellular region of a class II MHC molecule or at least part
of the extracellular region of the class II MHC molecule having an amino

acid sequence comprising one or more deletions, substitutions or additions. The molecule of the invention may be useful for detecting an antigen-specific CD4+ T-cell by flow cytometry and for presenting a microorganism-derived mucous membrane invasive protein as an antigen. The method of the invention enables efficient detection of antigen-specific activation of CD4+ T-cells in the mucous membrane. The current sequence is that of the class II major histocompatibility complex-related I-Ab(alpha)-Cholera toxin B subunit (CTB)-leucine zipper (LZ)-BirA fusion cDNA of the invention.

Sequence 945 BP; 230 A; 256 C; 294 G; 165 T; 0 U; 0 Other;

ery Match 69.9%; Score 665.2; DB 12; Length 945;
st Local Similarity 86.6%; Pred. No. 3.8e-145;
tches 759; Conservative 0; Mismatches 108; Indels 9; Gaps 2;

63 GTGTCGTGATGGTGTGTGACACGCCCGGGACTGAGGGCGGAAATCCATCTGCTTCGTCG 122
|||||
49 GTGACACTGATGGTGTGAGCTCCCACTGGCTTTTGGCTGGAGACTCTCGCGTGGGAAC 108
|||||
123 CCGTCGTGAGACACCGATCTGTGTGTCCGGCAGCTGGGACGAGAGTGGGGGCTCACTA 182
109 AATGAAGACGGCCACGGCATCGGGCCATCAGCATGGCGAA CGGAGGTGGTGGGTCCGGT 168
183 GTGCCCGGAGGCTCTGGAGTGGAGGTCGGAAGGCATTTCTGTGTCCAGTTCAAGGGC 242
169 GGAGGGGGAAG---TGGAGGTGGAGGCTCTGAAAGGCATTTCTGTGTACAGTTTCATGGC 225
243 GAGTGTCTACTACCAACGGGACGCGAGCGCATAGGCTCTGTGACCGATACATCTACCAAC 302
226 GAGTGTCTACTTCAACAA CGGGACGCGCATACGATATGTGACCAGATACATCTACCAAC 285
303 CGGAGAGGTACGTGCGCTACGACAGCGAGCTGGGCGAGTACCGCGGTGACCGAGCTG 362
286 CGGAGAGGTACGTGCGCTACGACAGCGAGCTGGGCGAGCACCGCGCGGTGACCGAGCTG 345
363 GGGCGGCGACGCGCGAGTACTTGGAAACAGCAGCAGCGGAGATCTCTGGAGCGAAACGGGGCC 422
346 GGGCGGCGACGCGCGAGTACTTGGAAACAGCAGCAGCGGAGATCTCTGGAGCGAAACGGGGCC 405
423 GAGTGTGACACGGGCTGTGACACAACTACGAGGGGCGGAGACCGACCTCTCTGGGG 482
406 GAGCTGTGACACGGTGTGACAGACAACCTACGAGGGGCGGAGACCCACACTCTCTCTGGGG 465
483 CGGCTTGAACAGGCCAATGTCCGCTCTGTCAGGACAGAGGCCCTCAACACCAAC 542
466 CGGCTTGAACAGGCCAATGTCCGCTCTGTCAGGACAGAGGCCCTCAACACCAAC 525
543 AACACTCTGGTCTGTTCCGTTGACAGATTTTACCCAGCCAAAGATCAAAGTGGCTGGTTC 602
526 AACACTCTGGTCTGCTCAGTGA CAGATTTTCTACCCAGCCAAAGATCAAAGTGGCTGGTTC 585
603 AGGAATGGCAGGAGGAGACAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGGAC 662
586 CGGAATGGCAGGAGGAGACGGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGGAC 645
663 TGGACCTTCCAGTCTCTGGTTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACC 722
646 TGGACCTTCCAGTCTCTGGTTCATGCTGGAGATGACCCCTCGGCGGGGAGAGGTCTACACC 705
723 TGGCATGTGGAGCATCCAGCCTGAAAGAGGCCCATCACTGTGGAGTGGAGGGGACAGATCC 782
706 TGTCACTGTGAGCATCCAGCCTGAGAGGCCCATCACTGTGGAGTGGAGGGGACAGATCG 765
783 GAGTCTCCCGGAGCAAGGAGGTGGAGGATCCACTACAGCTCCATCATGCTCAGTGTGAAA 842
766 TCAGCAGACC-----TGGTTCCGCGCGGATCCACTACAGCTCCATCAGTCACTGTGAAA 819
843 AAGAAATTTGCAACGACTGAGAAAGAAAGACGCTCAGCTGAATGGGAACTTCAAGCCCTC 902
820 AAGAAATTCGAGGCACTTAAGAAAGAAAGACGCTCAGCTGAATGGGAACTTCAAGCCCTC 879
903 AAGAAAGAAATTCGCGCCAGCATCATCATCATCAT 938

Db	880	AAAGAAGAACTGCCCGAGCTGCATCATATTCTGGAT	915
RESULT 4			
AAT04269			
ID	AAT04269	standard; DNA; 1013 BP.	
XX			
AC	AAT04269;		
XX			
DT	16-APR-1996	(first entry)	
XX			
DE	Hybrid IA beta chain gene.		
XX			
KW	Major histocompatibility complex; MHC; T-cell receptor; TCR;		
KW	autoimmune disease; immunodeficiency disease; immune response;		
KW	immunoproliferation disease; graft-host rejection; therapy; B cell;		
KW	M12.C3; pM12-IAB-Ea; SS.		
XX			
OS	Synthetic.		
XX			
FH	Key	Location/Qualifiers	
primer_bind	1..18		
FT	/*tag= a		
FT	/note= "probable primer binding site (primer #76)"		
primer_bind	complement(40..74)		
FT	/*tag= b		
FT	/note= "binding site for primer #362 (see AAT04270)"		
CDS	63..959		
FT	/*tag= c		
FT	/product= "hybrid IA beta chain"		
FT	63..143		
sig_peptide	/*tag= d		
FT	/note= "leader region"		
primer_bind	complement(140..191)		
FT	/*tag= e		
FT	/note= "binding site for primer #363 (see AAT04271)"		
primer_bind	complement(177..226)		
FT	/*tag= f		
FT	/note= "primer #364 binding site"		
primer_bind	complement(212..266)		
FT	/*tag= g		
FT	/note= "primer #365 (see AAT04272) binding site"		
primer_bind	385..403		
FT	/*tag= h		
FT	/note= "probable primer binding site (primer #270)"		
mat_peptide	531..959		
FT	/*tag= i		
FT	/product= "IA beta chain beta 2 region"		
primer_bind	535..564		
FT	/*tag= j		
FT	/note= "probable primer binding site (primer #271)"		
primer_bind	544..568		
FT	/*tag= k		
FT	/note= "probable primer binding site (primer #272)"		
primer_bind	823..850		
FT	/*tag= l		
FT	/note= "probable primer binding site (primer #259)"		
primer_bind	942..976		
FT	/*tag= m		
FT	/note= "probable primer binding site (primer #366)"		
primer_bind	1000..1013		
FT	/*tag= n		
FT	/note= "probable primer binding site (primer #59)"		
XX			
PN	WO9523814-A1.		
XX			
PD	08-SEP-1995.		
XX			
PF	03-MAR-1995;	95WO-US002689.	
XX			
PR	04-MAR-1994;	94US-00207481.	
XX			

Example 17; Page 137-139; 217pp; English.

The present sequence was used in the construction of major histocompatibility complex (MHC) fusion complexes

Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

ery Match 63.9%; Score 607.8; DB 2; Length 1508;
st Local Similarity 89.6%; Pred. No. 1.1e-131;
tches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;

21 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGTGCTG 80
6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGTGCTG 65
81 AGCAGCCCGGAGCTGAGGGCGGAATCCATCTGCTTCTGCCCTGCTGGAGCACCGG 140
66 AGCAGCCCAAGGAC-----CTTAAGTATCTCTCAGGCTGTTTAC 104
141 ATCGTGGTGTCCGCGAGCTGGAGCGAGTGGGGCTCACTAGTGCCTCCGAGGCTCTGGA 200
105 GCTGCTCAGCTGAAATCAACGAAGCTGTGTGTAGCGGAGGGGGCGGAAGCGCGGA 164
201 GGTGGAGGCTCCGAAAGGCATTTCTGTGTCCAGTTCAAGGGCGAGTCTACTACACCAAC 260
165 GGGGGAAGCTCCGAAAGGCATTTCTGTGTCCAGTTCAAGGGCGAGTCTACTACACCAAC 224
261 GGGAGCGAGATACGCTGTGACCAAGATACATCTACACCAAGAGTACGTGCGC 320
225 GGGAGCGAGATACGCTGTGACCAAGATACATCTACACCAAGAGTACGTGCGC 284
321 TAGCAGAGGAGCTGGGGGAGTACCGCGGTGACCGAGTGGGGCGCCAGACCGCGAG 380
285 TAGCAGAGGAGCTGGGGGAGTACCGCGGTGACCGAGTGGGGCGCCAGACCGCGAG 344
381 TACTGGAAACAGCCAGCCGAGATCTTGGAGCGAAGCGCGGGCGAGTGGACACCGCGTGC 440
345 TACTGGAAACAGCCAGCCGAGATCTTGGAGCGAAGCGCGGGCGAGTGGACACCGCGTGC 404
441 AGACACAATAGCAGGGGGCGAGACACGACCTCTCTCGGGCGGCTTGAACAGCCCAAT 500
405 AGACACAATAGCAGGGGGCGAGACACGACCTCTCTCGGGCGGCTTGAACAGCCCAAT 464
501 GTCGGCATCTCCCTGTCCAGGACAGAGCGCCCTCAACCAACCACTCTGTGTCTGTCG 560
465 GTCGGCATCTCCCTGTCCAGGACAGAGGGCCCTCAACCAACCACTCTGTGTCTGTCG 524
561 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGCCTGTTTCAAGAAATGGCCAGGAGAG 620
525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGCCTGTTTCAAGAAATGGCCAGGAGAG 584
621 ACAGTGGGGGTCTATCCACACAGCTTATTAGGAATGGGAGTGGACCTTCCAGTCTCTG 680
585 ACAGTGGGGGTCTATCCACACAGCTTATTAGGAATGGGAGTGGACCTTCCAGTCTCTG 644
681 GTCATCTGGAGATGACCCCTCATCAGGGAGAGGTCTACACCTGTCATGTGGAGATCCC 740
645 GTCATCTGGAGATGACCCCTCATCAGGGAGAGGTCTACACCTGTCATGTGGAGATCCC 704
741 AGCCTGAAGAGCCCCATCACTGTGAGTGA 771
705 AGCCTGAAGAGCCCCATCACTGTGAGTGA 735

LT 11
9069

AAx89069 standard; DNA; 1508 BP.

AAx89069;

14-SEP-1999 (first entry)

DE Single chain IAD/OVA 323-229 MHC fusion protein encoding DNA.
XX Major histocompatibility complex; MHC; single chain MHC; sc-MHC; Ig;
KW peptide binding groove; immunoglobulin; T cell receptor; immune response;
KW immune-related disorder; antigenic peptide; fusion protein; aa.
XX Synthetic.
OS WO9921572-A1.
XX 06-MAY-1999.
PD 13-OCT-1998; 98WO-US021520.
XX 29-OCT-1997; 97US-00960190.
XX (SUNO-) SUNOL MOLECULAR CORP.
XX Rhode PR, Acevedo J, Burkhardt M, Jiao J, Wong HC;
PI WPI; 1999-418411/35.
XX P-PSDB; AAY27111.
XX Single chain major histocompatibility complex class I complexes.
PT Example 1; Fig 1; 148pp; English.

XX The invention relates to new single chain major histocompatibility
CC complex (sc-MHC) class II complexes that comprise a peptide binding
CC groove, and a modified class II beta 2 chain or covalently linked
CC immunoglobulin (Ig) light chain constant (CI) region. The MHC complexes
CC are useful for detection and analysis of peptide ligands, pathogenic T-
CC cells, for functional, cellular and molecular assays. They can be used to
CC identify and isolate T cell receptor and/or MHC agonists and antagonists.
CC They can be used in vivo to compete with pathogenic antigen presenting
CC cells involved in immune-related disorders. They can also be used to
CC raise antibodies and to screen immune cells. It is also use in a method
CC of suppressing an immune response in mammals. The sc-MHC complexes
CC comprising modified class II beta 2 chains and/or Ig-CI regions are
CC soluble and provide enhanced yield. These MHC complexes also can contain
CC single antigenic peptides readily isolated from expressing cells in
CC significant quantities. The polypeptide MHC complexes also provide a
CC means to detect cells expressing multiple target structures with a single
CC complex. The present sequence represents a DNA encoding a single chain
CC IAD/OVA 323-229 MHC fusion protein
XX

SQ Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

Query Match 63.9%; Score 607.8; DB 2; Length 1508;
Best Local Similarity 89.6%; Pred. No. 1.1e-131;
Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;

QY 21 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGTGCTG 80
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGTGCTG 65
QY 81 AGCAGCCCGGAGCTGAGGGCGGAATCCATCTGCTTCTGCCCTGCTGGAGCACCGG 140
Db 66 AGCAGCCCAAGGAC-----CTTAAGTATCTCTCAGGCTGTTTAC 104
QY 141 ATCGTGGTGTCCGCGAGCTGGAGCGAGTGGGGCTCACTAGTGCCTCCGAGGCTCTGGA 200
Db 105 GCTGCTCAGCTGAAATCAACGAAGCTGTGTGTAGCGGAGGGGGCGGAAGCGCGGA 164
QY 201 GGTGGAGGCTCCGAAAGGCATTTCTGTGTCCAGTTCAAGGGCGAGTGTCTACTACACCAAC 260
Db 165 GGGGGAAGCTCCGAAAGGCATTTCTGTGTCCAGTTCAAGGGCGAGTGTCTACTACACCAAC 224
QY 261 GGGAGCGAGGATACGCTGTGACCAAGATACATCTACACCAAGAGAGTACGTGCGC 320
Db 225 GGGAGCGAGGATACGCTGTGACCAAGATACATCTACACCAAGAGAGTACGTGCGC 284
QY 321 TACGACAGGAGCTGGGGGAGTACCGCGGTGACCGGAGTGGGGCGCCAGACCGCGAG 380

105 GCTGCTCAGCTGAAATCAAGCAAGCTGCTGCTAGCGGAGGGCGGAAGCGCGGA 164
201 GGTGAGGCTCGAAGGCAATTTCTGTGTCAGATTCAAGGGCGAGTCTACTACACCAAC 260
165 GGGGGAAATCCGAAGGCAATTTCTGTGTCAGATTCAAGGGCGAGTCTACTACACCAAC 224
261 GGGAGCGCAGCGCATACGGCTCTGTGACCAAGATACATCTACAAACCGGAGGAGTACCTGGCG 320
225 GGGAGCGCAGCGCATACGGCTCTGTGACCAAGATACATCTACAAACCGGAGGAGTACCTGGCG 284
321 TACGACGACGACGTCGGCGAGTACCGCGGTGACCGAGCTGGCGCGCGCAGACGCCGAG 380
285 TACGACGACGACGTCGGCGAGTACCGCGGTGACCGAGCTGGCGCGCGCAGACGCCGAG 344
381 TACTGGACACGACCGGAGATCTGGAGGAAACGCGGCGGAGGTGGACACGCGCTGC 440
345 TACTGGACACGACCGGAGATCTGGAGGAAACGCGGCGGAGGTGGACACGCGCTGC 404
441 AGACACAACTACGAGGGCGGAGACCAAGCAGCTCTCTGCGGCGGCTTGAACAGCCCAAT 500
405 AGACACAACTACGAGGGCGGAGACCAAGCAGCTCTCTGCGGCGGCTTGAACAGCCCAAT 464
501 GTGCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCAACCAACACTCTGCTGTTCG 560
465 GTGCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCAACCAACACTCTGCTGTTCG 524
561 GTGACAGATTTTACCCAGCCAGATCAAGTGCCTGTTTCAGGAATGGCCAGGAGGAG 620
525 GTGACAGATTTTACCCAGCCAGATCAAGTGCCTGTTTCAGGAATGGCCAGGAGGAG 584
621 ACAGTGGGGGTCTATCCACACAGCTTATTAGGAATGGGGAATGGACCTTCCAGGTCTCTG 680
585 ACAGTGGGGGTCTATCCACACAGCTTATTAGGAATGGGGAATGGACCTTCCAGGTCTCTG 644
681 GTCATGCTGGAGATACCCCTCATCAGGAGAGGTTTACACCTGCGCATGTGGAGATCCC 740
645 GTCATGCTGGAGATACCCCTCATCAGGAGAGGTTTACACCTGCGCATGTGGAGATCCC 704
741 AGCCTGAGAGCCCCATCACTGTGAGTGA 771
705 AGCCTGAGAGCCCCATCACTGTGAGTGA 735

LT 15
7587
AAT17587 standard; DNA; 1508 BP.
AAT17587;
26-SEP-1996 (first entry)
Vector SCT1-derived single chain gene encoding MHC fusion complex.
MHC; major histocompatibility complex; PCR; polymerase chain reaction;
T cell activity modulator; antagonist; immune disorder; allergy;
multiple sclerosis; insulin-dependent diabetes mellitus;
rheumatoid arthritis; myasthenia gravis; ds.
Synthetic.
Key Location/Qualifiers
CDS 6..1508
/*tag= a
sig_peptide 6..86
/*tag= b
/*label= I-Ad beta chain leader
/*notes= "murine MHC class II I-Ad gene beta chain leader
sequence"
misc_feature 87..137
/*tag= c
/*label= OVA_323-339
/*note= "chicken ovalbumin residues 323-339"
138..167
/*tag= d
/*note= "10 residue linker peptide"
168..452
/*tag= e
/*label= I-Ad beta1
/*note= "murine MHC class II I-Ad gene beta-1 domain"
453..734
/*tag= f
/*label= I-Ad beta2
/*note= "murine MHC class II I-Ad gene beta-2 domain"
735..806
/*tag= g
/*note= "24 residue peptide linker"
807..1067
/*tag= h
/*label= I-Ad alpha1
/*note= "murine MHC class II I-Ad gene alpha-2 domain"
1068..1352
/*tag= i
/*label= I-Ad alpha2
/*note= "murine MHC class II I-Ad gene alpha-2 domain"
1353..1505
/*tag= j
/*label= I-Ad alpha-TM
/*note= "murine MHC class II I-Ad gene alpha-transmembrane
domain"
WO9604314-A1.
15-FEB-1996.
31-JUL-1995; 95WO-US009816.
29-JUL-1994; 94US-00283302.
01-FEB-1995; 95US-00382454.
XX (DADE-) DADE INT INC.
XX Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
PI Chavallaz P, Jiao J;
XX WPI; 1996-129343/13.
DR P-PSDB; AAR98906.
XX Major histocompatibility complex fusion complex for modulating T cell
PT activity - used in the treatment of immune disorders, e.g. multiple
PT sclerosis, IDDM and rheumatoid arthritis.
XX Example 17; Fig 28; 210pp; English.
XX AAT17587 encodes a murine MHC fusion complex capable of modulating T cell
CC activity encoded by the vector SCT1. The MHC fusion complex comprises at
CC least one MHC molecule containing a peptide-binding groove and a
CC presenting peptide covalently linked to the MHC molecule and opt. a
CC transmembrane domain. DNA encoding a MHC fusion complex may be cloned
CC into a host cell to express the complex. The transformed cells may then
CC be used to identify peptides that modulate, pref. antagonise, T cell
CC activity. DNA encoding a MHC fusion complex or a single chain fusion
CC molecule may be used to vaccinate a mammal against a targeted disorder.
CC The fusion complexes may be used to suppress an immune response in an
CC animal suffering from an immune disorder e.g. multiple sclerosis, insulin
CC -dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or
CC chronic allergies. The complexes may also be used in the treatment of
CC livestock and pets such as cats and dogs. The MHC fusion complexes can be
CC produced such that they contain a single antigenic peptide including one
CC of known structure, additionally a wide range of peptides can be
CC presented for T cell interaction
XX Sequence 1508 BP; 337 A; 414 C; 440 G; 317 T; 0 U; 0 Other;
SQ Query Match 63.7%; Score 606.2; DB 2; Length 1508;
Best Local Similarity 89.5%; Pred. No. 2.5e-131;

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protein - nucleic search, using frame_plus_p2n model
on: May 31, 2006, 23:04:55 / Search time 714 Seconds
(without alignment)
4482.160 Million cell updates/sec

US-10-048-116B-6
ect score: 1620
ence: 1 MALQIPSLLSNAVVVLVL.....LKKWLQALKKLAQHHEHHH 306

ing table: BLOSUM62

Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

ched: 5244920 seqs, 3486124231 residues

l number of hits satisfying chosen parameters: 10489840

um DB seq length: 0

um DB seq length: 2000000000

rocessing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

and line parameters:

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'N Geneseq -OFMT=fastap -SUFFIX=p2n.rng -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
'NS-bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
'ALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
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'N TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
'POP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

ibase : N Geneseq 8.*

- 1: Geneseqn1980s.*
- 2: Geneseqn1990s.*
- 3: Geneseqn2000s.*
- 4: Geneseqn2001as.*
- 5: Geneseqn2001bs.*
- 6: Geneseqn2002as.*
- 7: Geneseqn2002bs.*
- 8: Geneseqn2003as.*
- 9: Geneseqn2003bs.*
- 10: Geneseqn2003cs.*
- 11: Geneseqn2003ds.*
- 12: Geneseqn2004as.*
- 13: Geneseqn2004bs.*
- 14: Geneseqn2005s.*
- 15: Geneseqn2006s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Alt	Score	Query	Length	ID	Description
1	1620	100.0	921	5 AAF55099	Aaf55099 DNA encod
2	1255.5	77.5	893	2 AAT04262	Aat04262 Hybrid IA
3	1235	76.2	945	12 ADQ31225	Adq31225 I-Ab(beta

4	1219	75.2	915	12	ADQ31228	Adq31228 I-Ab(beta
5	1161.5	71.7	4724	2	AAV12068	Aav12068 Murine IA
6	1151	71.0	1013	2	AAT04269	Aat04269 Hybrid IA
7	1145	70.7	1382	2	AAT17588	Aat17588 Vector SC
8	1145	70.7	1382	2	AAT86989	Aat86989 SCE1 sing
9	1145	70.7	1382	8	ACA60744	Acac60744 Mouse MHC
10	1145	70.7	1385	2	AAT17586	Aat17586 Vector SS
11	1145	70.7	1385	2	AAT86987	Aat86987 SSC1 sing
12	1145	70.7	1385	8	ACA60742	Acac60742 Mouse MHC
13	1145	70.7	1508	2	AAT17587	Aat17587 Vector SC
14	1145	70.7	1508	2	AAT86988	Aat86988 SCT1 sing
15	1145	70.7	1508	2	AAX89069	Aax89069 Single ch
16	1145	70.7	1508	8	ACA60743	Acac60743 Mouse MHC
17	1096.5	67.7	798	12	ADJ75986	Adj75986 Marker ge
18	1096.5	67.7	798	14	ADX26090	Adx26090 Novel cel
19	1093	67.5	1085	4	ABI99040	Abi99040 Murine PC
20	1049.5	64.8	1698	4	ABI99038	Abi99038 Murine PC
21	1044.5	64.5	1662	4	ABI99039	Abi99039 Murine PC
22	979.5	60.5	702	2	AAQ03170	AAq03170 Sequence
23	979.5	60.5	702	2	AAT06286	Aat06286 I-Ab-beta
24	979.5	60.5	702	2	AAQ56920	AAq56920 Mouse I-A
25	972	60.0	1243	6	ABN84048	Abn84048 Single ch
26	963.5	59.5	702	2	AAQ35055	AAq35055 TAB beta
27	957	59.1	1686	4	ABI99031	Abi99031 MBP 1-14
28	957	59.1	1701	4	ABI99028	Abi99028 TAS MBP 1
29	957	59.1	2059	4	ABI99032	Abi99032 MBP 1-14
30	957	59.1	2346	4	ABI99027	Abi99027 TAS MBP 1
31	952	58.8	1707	4	ABI99030	Abi99030 TAS MBP 9
32	949	58.6	1680	4	ABI99021	Abi99021 I-A8 MBP
33	949	58.6	2053	4	ABI99029	Abi99029 TAS MBP 9
34	949	58.6	2343	4	ABI99033	Abi99033 MBP 90-10
35	871	53.8	1344	2	AAT60705	Aat60705 cDNA enco
36	854.5	52.7	1323	2	AAT60700	Aat60700 cDNA enco
37	844.5	52.1	861	14	ASC64482	Aac64482 DRB1-biot
38	839.5	51.8	1192	10	AAQ63150	AAq63150 Human maj
39	839.5	51.8	1192	10	AAQ62751	AAq62751 Human maj
40	839.5	51.8	1192	11	ADP88246	Adp88246 Lung canc
41	839.5	51.8	1192	13	ADR24869	Adr24869 Breast ca
42	833.5	51.5	941	12	ADO40822	Ado40822 DNA encod
43	829	51.2	562	6	ABK63510	Abk63510 Rat seque
44	829	51.2	562	10	ADB57995	Adb57995 Toxicity-
45	829	51.2	562	10	ABT41775	Abt41775 Toxicity

ALIGNMENTS

RESULT 1

AAF55099

ID AAF55099 standard; DNA; 921 BP.

XX AAF55099;

DT 15-MAY-2001 (first entry)

DE DNA encoding a fusion protein comprising a beta chain of MHC.

XX Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
KW major histocompatibility complex; Fc region; antigen; T lymphocyte;
KW immunostimulant; vaccine; infection; tumour; ss.

XX Synthetic.

XX Key Location/Qualifiers

FT CDS 1..921

FT /*tag= a

XX WO200109194-A1.

XX 08-FEB-2001.

XX 28-JUL-2000; 2000WO-FR002193.

XX 29-JUL-1999; 99FR-00009862.

(CNRS) CNRS CENT NAT RECH SCI.

Glaichenhaus N, Malherbe L;

WPI; 2001-182944/18.

P-PSDB; AAB67481.

New soluble recombinant protein, useful e.g. as immunostimulant, comprises dimeric major histocompatibility complex molecule fused to immunoglobulin Fc region.

Example 1: page 34-35; 43pp; French.

The specification describes soluble recombinant proteins that comprise at least a dimer formed from the alpha and beta-chains of MHC (major histocompatibility complex) Class I and II molecules in which at least one chain has, attached to its C-terminus, at least part of the Fc region of an immunoglobulin. The recombinant proteins, when linked to an antigenic peptide, are used to count and/or purify antigen-reactive T lymphocytes and to characterize their phenotype, e.g. in preclinical evaluation of vaccines. They are also used as immunostimulants, particularly for vaccine development (against infections and tumours), to count and determine phenotype of autoreactive T cells in subjects with, or at risk of developing, autoimmune diseases, e.g. for staging or evaluating treatments, and (to purify and/or enrich Ag-reactive T cells from cell cultures or patient samples, for use in subsequent curative or preventative cellular therapy. The present sequence encodes a recombinant protein of the invention, comprising a beta chain of MHC molecules

Sequence 921 BP; 214 A; 265 C; 286 G; 156 T; 0 U; 0 Other;

ment Scores:		
a. NO.:	6.68e-144	921
ent Similarity:	1620.00	306
Local Similarity:	100.0%	Conservative: 0
y Match:	100.0%	Mismatches: 0
	5	Indels: 0
		Gaps: 0

J-048-116B-6 (1-306) x AAF55099 (1-921)

1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
|||||
1 ATGGCTCTGCAGATCCCGACGCTCTCTCTCAGCTGCCTGCTGCTGCTGCTGCTG 60
|||||
21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
|||||
61 AGCAGCCCCGGGACTGAGGGCGGAAATCCATCTGCTTCTGCGCTGCTGAGACACCCG 120
|||||
41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
|||||
121 ATCGTGGTGTCCGGCAGCTGGGACGGAGGTGGGGGCTCACTAGTGGCCCCGAGGCTCTGGA 180
|||||
61 GlyGlyGlySerGluArgHisPheValValGlnPheIysGlyGlyCysTyrTrpThrAsn 80
|||||
181 GGTGGAGGCTCCGAAAGGCAATTTCTGGTGCAGTTTCARGGCGGAGTGTACTACACCAAC 240
|||||
81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 300
|||||
241 GGGACGCGGGGCATACGGCTCGTGACCAAGATACATCTACACCGGAGGAGTACGTGGCG 360
|||||
101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
|||||
301 TACGACAGCGACCTGGGCGGAGTACC CGCGGTACCGAGCTGGGGCGGCGACAGCGCCGAG 360
|||||
121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
|||||
361 TACTGGAAACGCCGCGGAGATCTCTGGACGAAACCGGGCCCGAGGTGGACACGGCGTCC 420
|||||
141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
|||||
421 AGACACAACTACGAGGGCGGAGACGAGCACTCTCTCGGCGGTCTGAACAGGCCCAAT 480

03-DEC-2002; 2002JP-00351818.

(SENT-) SENTAN KAGAKU GIJUTSU INCUBATION CENT KK.

WPI; 2004-546819/53.
P-PSDB; ADQ31224.

Peptide-Class II major histocompatibility complex (MHC) composite, useful for detecting antigen specific CD4+ T-cell, comprises antigen peptide containing epitope of mucous membrane invasive protein, and extracellular region of MHC.

Example 1; SEQ ID NO 10; 30pp; Japanese.

The invention relates to a novel class II major histocompatibility complex (MHC) antigenic peptide composite comprising a peptide containing the T-cell antigenic determinant of a mucous membrane invasive protein and the extracellular region of a class II MHC molecule or at least part of the extracellular region of the class II MHC molecule having an amino acid sequence comprising one or more deletions, substitutions or additions. The molecule of the invention may be useful for detecting an antigen-specific CD4+ T-cell by flow cytometry and for presenting a microorganism-derived mucous membrane invasive protein as an antigen. The method of the invention enables efficient detection of antigen-specific activation of CD4+ T-cells in the mucous membrane. The current sequence is that of the class II major histocompatibility complex-related I-Ab(alpha)-Cholera toxin B subunit (CTB)-leucine zipper (LZ)-Bira fusion cDNA of the invention.

Sequence 945 BP; 230 A; 256 C; 294 G; 165 T; 0 U; 0 Other;

ment Scores:

No.:	2, 27e-107	Length:	945
e:	1235.00	Matches:	244
ent Similarity:	84.9%	Conservative:	14
Local Similarity:	80.3%	Mismatches:	42
y Match:	76.2%	Indels:	4
	12	Gaps:	3

3-048-116B-6 (1-306) x ADQ31225 (1-945)

```

1 MetAlaLeuGlnIlePro---SerLeuLeuLeuSerAlaAlaValValLeuMetVal 19
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
4 GTGTCTGAGCTCCCTGGAGTCTCTACATGCGCAAGCTGACAGTGCACACTGATGTG 63

20 LeuSerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHis 39
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
64 CTGAGCTCCCACTGGCTTTGGCTGGAGACTCTCTCGTGTGGAAACAATAAGACGCGCAC 123

40 ProfileValValSerGlySerTrpAspGlyGlyGlyGlySerLeuValProArgGlySer 59
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
124 GCGATCGCGCCCATCAGCATGCGCAACGGAGGTGTGGTCC---GGTGGAGGGGAAGT 180

60 GlyGlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTrpThr 79
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
181 CGAGGTGGAGGGTCTGMAAGGCATTCGTGTACCAAGTTTCATGGCGAGTGTACTTACC 240

80 AsnGlyThrGlnArgIleArgLeuValThrArgTrpIleTrpAsnArgGluGluTrpVal 99
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
241 AACGGGACGCGCATACGATATGTGACCGATCATCTACACCGGGAGGAGTACGTG 300

100 ArgTrpAspSerAspValGlyGlyTrpArgAlaValThrGluLeuGlyArgProAspAla 119
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
301 CGCTACGACAGCGAGTGGCGGAGCAGCCGGCGGTGACCGAGCTGGGGGGCCAGACGCC 360

120 GluTrpTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAla 139
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
361 GAGTACTGGAAACAGCCAGCCGAGATCTCTGGAGCGAAGCGGGCGCGTGGACACGGTG 420

140 CysArgHisAsnTrpGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnPro 159
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
421 TGCAGACACAACTACGAGGGCGCGGAGACCCACACTCTCTCGCGCGCTTGAAACGCC 480

```

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QY 160 AsnValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCys 179
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 481 AATGTGGTCATCTCCCTGTCCAGGACAGAGCCCTCAACACCACACACACTCTGGTCTGC 540

QY 180 SerValThrAspPheTrpProAlaIleLysValArgTrpPheArgAsnGlyGlnGlu 199
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 541 TCAGTGCAGAGATTTCTACCCAGCAAGATCAAGTGCCTGTTCCGGAATGGCCAGGAG 600

QY 200 GluThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnVal 219
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 601 GAGACGGTGGGGTCTCATCCACAGCTTATTAGAATGGGACTGGACCTTCCAGGTC 660

QY 220 LeuValMetLeuGluMetThrProHisGlnGlyGluValTrpCysHisValGluHis 239
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 661 CTGGTCATGTGGAGATGACCCCTCGGGGGGAGAGGTCTACACCTGTCACTGGAGCAT 720

QY 240 ProSerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSer 259
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 721 CCCAGCCTGAAGAGCCCATCATCTGTGGAGTGGAGGCGACAGTCTGTACGCA-----GAC 774

QY 260 LysGlyGlyGlySerThrAlaProSerAlaGlnLeuLysValLysLeuGlnAla 279
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 775 CTGGTTCGCGCGGATCCACTCAGCTCCATCAGCTCAGTTGMAAAGAAAGAACTGCAGGCA 834

QY 280 LeuLysLysLysAsnAlaGlnLeuLysTrpLysLeuGlnAlaLeuLysLysLeuAla 299
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 835 CTTAAGAAAGAAAGACGCTCAGCTGAAGTGGAAACTTCAAGCCCTCAAGAAGAAACTCGCC 894

QY 300 GlnHisHisHis 303
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 895 CAGCTGCATCAT 906

```

RESULT 4

ADQ31228
ID ADQ31228 standard; cDNA; 915 BP.

AC ADQ31228;

XX 07-OCT-2004 (first entry)

XX I-Ab(beta)-E. coli heat-labile toxin B subunit-LZ-Bira fusion cDNA.

XX class II major histocompatibility complex; MHC; CD4+ T-cell detection;

XX flow cytometry; mucous membrane invasive antigen;

XX I-Ab(beta)-heat-labile toxin B subunit-leucine zipper-Bira fusion; LTB;

XX ss; gene.

XX Escherichia coli.

OS Unidentified.

XX Key Location/Qualifiers

XX CDS 1..915

XX /start= a

XX /product= "I-Ab(beta)-Escherichia coli heat-labile toxin

XX B subunit (LTB)-leucine zipper (LZ)-Bira fusion protein"

XX JP2004196789-A.

XX 15-JUL-2004.

XX 03-DEC-2003; 2003JP-00404367.

XX 03-DEC-2002; 2002JP-00351818.

XX (SENT-) SENTAN KAGAKU GIJUTSU INCUBATION CENT KK.

XX WPI; 2004-546819/53.

XX P-PSDB; ADQ31227.

XX Peptide-Class II major histocompatibility complex (MHC) composite, useful for detecting antigen specific CD4+ T-cell, comprises antigen peptide containing epitope of mucous membrane invasive protein, and extracellular

region of MHC.

Example 3; SEQ ID NO 13; 30pp; Japanese.

The invention relates to a novel class II major histocompatibility complex (MHC) antigenic peptide composite comprising a peptide containing the T-cell antigenic determinant of a mucous membrane invasive protein and the extracellular region of a class II MHC molecule or at least part of the extracellular region of the class II MHC molecule having an amino acid sequence comprising one or more deletions, substitutions or additions. The molecule of the invention may be useful for detecting an antigen-specific CD4+ T-cell by flow cytometry and for presenting a microorganism-derived mucous membrane invasive protein as an antigen. The method of the invention enables efficient detection of antigen-specific activation of CD4+ T-cells in the mucous membrane. The current sequence is that of the class II major histocompatibility complex-related I-Ab(alpha)-Escherichia coli heat-labile toxin B subunit (LTB)-leucine zipper (LZ)-BirA fusion cDNA of the invention.

Sequence 915 BP; 228 A; 242 C; 271 G; 174 T; 0 U; 0 Other;

ment Scores:

i. No.: 7,14e-106 Length: 915
e: 1219.00 Matches: 242
ent Similarity: 84.9% Conservative: 16
Local Similarity: 79.6% Mismatches: 32
y Match: 75.2% Indels: 14
12 Gaps: 5

.0-048-116B-6 (1-306) x ADQ31228 (1-915)

1 MetAlaLeuGlnIlePro---SerLeuLeuSerAlaAlaValValLeuMetVal 19
4 GRGTGTCGAAGCTCCCTCGAGGTTCCTACATGGCAAGCTGACAGTGCACGTGAGTG 63
20 LeuSerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHis 39
64 CTGAGCTCCCACTGGCTTGGCTGGAGACTCC-----GGTAAAGAGAA 108
40 ProIleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySer 59
109 ATGGTTATCATTT-----ACATTTAAGGTGGTGGTGGTCTTTAGTTCTTACA 156
60 GlyGlyGlyGlySerGluArgHisPheValValGlnPheLeuGlyGlyCysTyrThr 79
157 -----GGTGGTAGTGAAGGCATTTCTGTACCATGTTCTGGCGGAGTGCTACTTACC 210
80 AsnGlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrVal 99
211 AACGGGACGCGGCATACATATGTGACCATATCTACACCGGAGGAGTACGTG 270
100 ArgTyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAla 119
271 CGTACGACAGCGACGTGGCGGAGCACCACCGCGGTGACCGAGCTGGCGGCGCAGACGCC 330
120 GluTyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaValAspThrAla 139
331 GAGTACTGGAACAGCGCGGAGATCTCGAGCGGAACCGCGCGGAGCTGACACAGGTG 390
140 CysArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluPro 159
391 TGCAGACACACTACAGCGGGCGGAGACCCACCTCTCTGGCGGGCTTGACAGCC 450
160 AsnValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCys 179
451 AATGTCGTATCTCCCTGTCCAGGACAGAGGCGCTCAACACCAACCACTCTGGTCTGC 510
180 SerValThrAspPheTyrProAlaTyrIleValArgTyrPheArgGlnGlu 199
511 TCAGTGACAGATTTCTTACCAGCGCAAGATCAAGTGGCTGGTTCGGAGTGGCCAGGAG 570
200 GluThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnVal 219

Db 571 GRAGCGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGAGCTTCCAGGTC 630
Qy 220 LeuValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHis 239
Db 631 CTGGTCATGCTGGAGATGACCCCTCGCGGGGAGAGGCTTACACCTGTCCAGTGGAGCAT 690
Qy 240 ProSerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSer 259
Db 691 CCCAGCTGTAGAGAGCCCATCACTGTGGAGTGGAGGACACAGTCTGTGTCACCA 744
Qy 260 LysGlyGlyGlySerThrAlaProSerAlaGlnLeuLysLysLysLeuGlnAla 279
Db 745 CTGGTTCGCGCGGATCCACTACAGCTCCATCAGCTCAGTTCGTAAGAGAACTGCAAGCA 804
Qy 280 LeuLysLysLysAsnAlaGlnLeuLysLysLysLeuGlnAlaLeuLysLysLysLeuAla 299
Db 805 CTTAAGAAAAAGAACGCTCAGCTGAAGTGGAAACTTCAAGCCCTCAAGAAGAACTCGCC 864
Qy 300 GlnHisHisHis 303
Db 865 CAGTGCATCAT 876
RESULT 5
AAV12068
ID AAV12068 standard; cDNA; 4724 BP.
XX
AC AAV12068;
XX
DT 08-JUN-1998 (first entry)
XX Murine IAD beta chain cDNA.
XX Major histocompatibility class II antigen; MHC class II; T cell;
KW T lymphocyte; Th; Th2; activation; CD4+; antigen presenting cell; APC;
KW autoimmune disease; diabetes; multiple sclerosis; autoimmune thyroiditis;
KW systemic lupus erythematosus; myasthenia gravis; Crohn's disease;
KW inflammatory bowel disease; allergy; asthma; contact sensitivity;
KW immunotherapy; therapy; IAD beta chain; mouse; da; circular; cyclic.
XX Mus musculus.
OS
XX WO9746256-A1.
PN
XX 11-DEC-1997.
PD
XX 22-MAY-1997; 97WO-US008697.
PF
XX 23-MAY-1996; 96US-0018175P.
PR
XX (SRI) SCHRIPPS RES INST.
PA
XX Webb SR, Wingvist O, Karlsson L, Jackson MR, Peterson PA;
PI WPI; 1998-041895/04.
DR
XX Synthetic antigen presenting cell for activating CD4+ T cells - useful to
PT treat autoimmune disease, e.g. diabetes, multiple sclerosis, Crohn's
PT disease and inflammatory bowel disease, or allergy, e.g. asthma and
PT contact sensitivity.
XX Example 2; Page 94-96; 141pp; English.
PS
XX This nucleotide sequence comprises a PCR product obtained by
CC amplification of mouse splenocyte cDNA using primers (see AAV12065 and
CC AAV12066) designed for the amplification of IAD beta chain full-length
CC cDNA. IAD alpha chain cDNA (see AAV12067) has been similarly obtained.
CC The IAD sequences were cloned into metallothionein promoter (see
CC AAV12062)-driven vector pMTa-3 prior to sequencing. Major
CC histocompatibility complex (MHC) class II IAD heterodimers were expressed
CC at the cell surface of transfected Drosophila Schneider 2 (ATCC CRL
CC 10974) cells. The invention relates to the preparation and use of
CC synthetic antigen presenting matrices, in particular antigen presenting
CC cells such as insect cells that have been transfected to produce MHC

antigen presenting molecules with one or more accessory molecules. The matrices are used to activate naive CD4+ T cells and to shift the ongoing activation state into a preferred differentiated population of Th1 or Th2 cells. Applications include the treatment of autoimmune disease, e.g. diabetes, multiple sclerosis, autoimmune thyroiditis, systemic lupus erythematosus, myasthenia gravis, Crohn's disease and inflammatory bowel disease, or an allergy, e.g. asthma and contact sensitivity

Sequence 4724 BP; 1196 A; 1194 C; 1200 G; 1134 T; 0 U; 0 Other;

ment Scores:

. No.:	1.85e-99	Length:	4724
3:	1161.50	Matches:	228
Int Similarity:	85.8%	Conservative:	1
Local Similarity:	85.4%	Mismatches:	1
/ Match:	71.7%	Indels:	37
	2	Gaps:	2

0-048-116B-6 (1-306) x AAV12068 (1-4724)

1	MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValMetValLeu	20
451	ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTATGTGTGTG	510
21	SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro	40
511	AGCAGCCCGGAGCTGAGGGCGGAAC	537
41	IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly	60
537	-----	537
61	GlyGlyGlySerGluArgHisPheValValGlnPheIleGlyCysTyrThrAsn	80
538	-----TCGAAGGCAATTCGTGTTCAGTTCAGGGCGAGTGTACTACCCAAAC	588
81	GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg	100
589	GGGAGCGAGCGCATACGGCTGTGACCATACATCTCAACCGGGAGGAGTACGTGGCG	648
101	TyrAspSerAspValGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu	120
649	TACGACGCGAGTGGGGAGTACCGCGGTGACCGAGCTGGGGCGCCAGACGCCGAG	708
121	TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys	140
709	TACTGGACAGCCAGCCGAGATCTCTGGAGCGAAGCGGGCGAGTGGACACGGCGTGC	768
141	ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn	160
769	AGACACAACCTACGAGGGCGGAGACCGACCTCCCTCGCGGGCTTGAACAGCCCAAT	828
161	ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer	180
829	ATCGCCATCTCCCTCTCAGGACAGAGGCCCTCAACCCACACACACTCTGTGTGTTCG	888
181	ValThrAspPheTyrProAlaIleIleValArgTyrPheArgAsnGlyGlnGluGlu	200
889	GTGACAGATTTCTACCCAGCCAGATCAAGTGCCTGTTTCAGGAATGGCCAGGAGAG	948
201	ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrThrPheGlnValLeu	220
949	ACAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCAGGTCTCG	1008
221	ValMetLeuGluMetThrProHisGlnGlyValValTyrThrCysHisValGluHisPro	240
1009	GTCTGCTGGAGATACCCCTCATCAGGAGAGGTCTACCTCTCATGTGGAGCATCC	1068
241	SerLeuIleSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerIle	260
1069	AGCCTGAAGAGCCCATCACTGTGAGTGGAGGCGACAGTCCGAGTCTGCCCGGAGCAG	1128
261	-----GlyGlyGlyGly	264

Db	1129	ATGTTGAGCGCATCGGGGC	1149
RESULT 6			
AAT04269			
ID	AAT04269	standard; DNA; 1013 BP.	
XX	AAT04269;		
XX	16-APR-1996	(first entry)	
XX	Hybrid IA beta chain gene.		
XX	Major histocompatibility complex; MHC; T-cell receptor; TCR;		
KW	autoimmune disease; immunodeficiency disease; immune response;		
KW	immunoproliferation disease; graft-host rejection; therapy; B cell;		
KW	M12.C3; pM12-IAB-Ea; 88.		
XX	Synthetic.		
XX	Key	Location/Qualifiers	
FT	primer_bind	1..18	
FT		/*tag= a	
FT	primer_bind	/note= "probable primer binding site (primer #76)"	
FT		complement(40..74)	
FT		/*tag= b	
FT		/note= "binding site for primer #362 (see AAT04270)"	
FT	CDS	63..959	
FT		/*tag= c	
FT	sig_peptide	/product= "hybrid IA beta chain"	
FT		63..143	
FT		/*tag= d	
FT	primer_bind	/note= "leader region"	
FT		complement(140..191)	
FT		/*tag= e	
FT	primer_bind	/note= "binding site for primer #363 (see AAT04271)"	
FT		complement(177..226)	
FT		/*tag= f	
FT	primer_bind	/note= "primer #364 binding site"	
FT		complement(212..266)	
FT		/*tag= g	
FT	primer_bind	/note= "primer #365 (see AAT04272) binding site"	
FT		385..403	
FT		/*tag= h	
FT	mat_peptide	/note= "probable primer binding site (primer #270)"	
FT		531..959	
FT		/*tag= i	
FT	primer_bind	/product= "IA beta chain beta 2 region"	
FT		535..564	
FT		/*tag= j	
FT	primer_bind	/note= "probable primer binding site (primer #271)"	
FT		544..568	
FT		/*tag= k	
FT	primer_bind	/note= "probable primer binding site (primer #272)"	
FT		823..850	
FT		/*tag= l	
FT	primer_bind	/note= "probable primer binding site (primer #259)"	
FT		942..976	
FT		/*tag= m	
FT	primer_bind	/note= "probable primer binding site (primer #366)"	
FT		1000..1013	
FT		/*tag= n	
FT		/note= "probable primer binding site (primer #59)"	
XX	WO9523814-A1.		
XX	08-SEP-1995.		
XX	03-MAR-1995;	95WO-US002689.	
XX	04-MAR-1994;	94US-00207481.	
XX	(NAJE-) NAT JEWISH CENT IMMUNOLOGY & RESPIRATORY.		

Kappler JW, Marrack P;

WPI; 1995-320543/41.
P-PSDB; AAR82538.

Peptide-MHC complex comprising antigenic peptide, linker and MHC segment - useful as reagents for the treatment of diseases including auto-immune diseases, immuno-stimulatory diseases or graft-host rejection.

Example 2; Page 65; 9app; English.

This sequence represents a hybrid IA beta chain gene. This sequence contains a fragment of the IE alpha chain (residues 56-73), as well as a linker and cleavage site. This sequence was transfected into a B cell line (M12.C3) using plasmid pM12-IAB-Ea. It was found that the encoded sequence was expressed in these cells. Complexes such as this may be used to regulate an immune response. The complexes are capable of being recognised by a TCR alone or in combination with additional MHC proteins. These complexes are useful for therapeutic purposes and experimental purposes. They can also be used as reagents for the treatment of diseases including autoimmune diseases, immunodeficiency diseases, immunoproliferation diseases, and graft-host rejection

Sequence 1013 BP; 220 A; 272 C; 327 G; 192 T; 0 U; 2 Other;

ment Scores:

i. No.: 2,31e-99 Length: 1013
e: 1151.00 Matches: 230
ent Similarity: 86.8% Conservative: 6
Local Similarity: 84.6% Mismatches: 22
y Match: 71.0% Indels: 14
Gaps: 2

10-048-116B-6 (1-306) x AAT04269 (1-1013)

1 MetAlaLeuGluIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20
63 ATGGCTCTCAGATCCCAAGCTTCCTCTGGCTGCTGGTGTGCTCATGGTGGCTG 122
21 SerSerProGlyThrGluGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
123 AGCAGCCAGGAGACTGAGCGCGGAGACTCC-----GAAGCTAGCTTGGAGGCTCAG 173
41 -----IleValValSerGlySerTrpAspGlyGlyGlySerLeuVal 55
174 GGTGCACTGCCCAACATTGCTGTCGACAAAGCTGGAGGTGGTGGATCCGGTGA----- 227
56 ProArgGlySerGlyGlyGlySerGluArgHisPheValValGlnPheGlyGlyGlu 75
228 ---GGGGGAAGTGGAGGTGGAGGTCTGAAGGCATTTCTGTATCCAGTTTCATGGCGGAG 284
76 CysTyrTyrThrAsnGlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArg 95
285 TGCTACTTCACCAACCGGAGCGAGCGCATACGATATGTGACCGACATACATCAACACCG 344
96 GluGluTyrValArgTyrAspSerAspValGlyGlyTyrArgAlaValThrGluLeuGly 115
345 GAGGAGTACGTGGCTACGACAGCAGTGGCGGCGAGCCGCGCGTGCACCGAGCTGGGG 404
116 ArgProAspAlaGluTyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGlu 135
405 CGGCCACAGCGCCAGTACTGGAAACACGCCCGGAGATCTGTGAGGAAACCGGCGCGGAG 464
136 ValAspThrAlaCysArgHisAenTyrGluGlyProGluThrSerThrSerLeuArgArg 155
465 GTGGACACGGTGTGCAGACACAACTACGAGGGGCCGAGACCCACACTCCCTGCGCGCGG 524
156 LeuGluGlnProAsnValAlaIleSerLeuSerArgThrGluAlaLeuAenHisHisAen 175
525 CTTGAACAGCCCAATGTCGTCTCTCCCTGTCCAGGACAGAGGCCCTCAACACCAACAAC 584
176 ThrLeuValCysSerValThrAspPheTyrProAlaLysIleLysValArgTrpPheArg 195

Db 585 ACTCTGGTCTGCTCAGTGACAGATTCTTACCCAGCCAGATCAAGTGGCTGGTCCGG 644
Qy 196 AenGlyGlnGluThrValGlyValSerSerThrGlnLeuIleArgAenGlyAspTrp 215
Db 645 AATGCCAGGAGAGACGGTGGGCTCTCATCCACACAGCTTATTAGGAATGGGACTGG 704
Qy 216 ThrPheGlnValLeuValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCys 235
Db 705 ACCTTCCAGGTCCTGCTCATGCTGAGATGACCCCTCGCGGGGAGAGGTCTATACCTGT 764
Qy 236 HisValGluHisProSerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGlu 255
Db 765 CACGTGGAGCATCCAGCTGAGAGCCCCCATCATCTGTGGAGTGGAGGGGCACAGTCTGAG 824
Qy 256 SerAlaArgSerLys-----GlyGlyGlyGly 264
Db 825 TCTGCTGGAGCAAGATGTTGAGCGGCATCGGGGGC 860
RESULT 7
AAT17588
ID AAT17588 standard; DNA; 1382 BP.
XX AAT17588;
DT 26-SEP-1996 (first entry)
XX Vector SCEI-derived single chain gene encoding MHC fusion complex.
DE MHC; major histocompatibility complex; PCR; polymerase chain reaction;
KW T cell activity modulator; antagonist; immune disorder; allergy;
KW multiple sclerosis; insulin-dependent diabetes mellitus;
KW rheumatoid arthritis; myasthenia gravis; ds.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT CDS 6..1382
FT /tag= a
FT sig_peptide 6..86
FT /tag= b
FT /label= I-Ad beta chain leader
FT /note= "murine MHC class II I-Ad gene beta chain leader
FT sequence"
FT misc_feature 87..137
FT /tag= c
FT /label= OVA 323-339
FT /note= "chicken ovalbumin residues 323-339"
FT misc_feature 138..167
FT /tag= d
FT /note= "10 residue linker peptide"
FT misc_feature 168..452
FT /tag= e
FT /label= I-Ad beta1
FT /note= "murine MHC class II I-Ad gene beta-1 domain"
FT misc_feature 453..734
FT /tag= f
FT /label= I-Ad beta2
FT /note= "murine MHC class II I-Ad gene beta-2 domain"
FT misc_feature 735..806
FT /tag= g
FT /note= "24 residue peptide linker"
FT misc_feature 807..1067
FT /tag= h
FT /label= I-Ad alpha1
FT /note= "murine MHC class II I-Ad gene alpha-1 domain"
FT misc_feature 1068..1352
FT /tag= i
FT /label= I-Ad alpha2
FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
FT misc_feature 1353..1379
FT /tag= j
FT /note= "EE tag"

WO9604314-A1.
15-FEB-1996.
31-JUL-1995; 95WO-US009816.
29-JUL-1994; 94US-00283302.
01-FEB-1995; 95US-00382454.
(DADE-) DADE INT INC.
Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
Chavallaz P, Jiao J;
WPI; 1996-129343/13.
P-PSDB; AAR98907.
Major histocompatibility complex fusion complex for modulating T cell
activity - used in the treatment of immune disorders, e.g. multiple
sclerosis, IDDM and rheumatoid arthritis.
Example 17; Fig 29; 210pp; English.
AAT17588 encodes a murine MHC fusion complex capable of modulating T cell
activity encoded by the vector SCE1. The MHC fusion complex comprises at
least one MHC molecule containing a peptide-binding groove and a
presenting peptide covalently linked to the MHC molecule and opt. a
transmembrane domain. DNA encoding a MHC fusion complex may be cloned
into a host cell to express the complex. The transformed cells may then
be used to identify peptides that modulate, pref. antagonise, T cell
activity. DNA encoding a MHC fusion complex or a single chain fusion
molecule may be used to vaccinate a mammal against a targeted disorder.
The fusion complexes may be used to suppress an immune response in an
animal suffering from an immune disorder e.g. multiple sclerosis, insulin
-dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or
chronic allergies. The complexes may also be used in the treatment of
livestock and pets such as cats and dogs. The MHC fusion complexes can be
produced such that they contain a single antigenic peptide including one
of known structure, additionally a wide range of peptides can be
presented for T cell interaction
Sequence 1382 BP; 320 A; 374 C; 404 G; 284 T; 0 U; 0 Other;
ment Scores:
No.: 1382
g: 1345.00
ant Similarity: 87.2%
Local Similarity: 85.7%
y Match: 70.7%
Indels: 10
Gaps: 3
J--048-116B-6 (1-306) x AAT17588 (1-1382)
1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
6 ATGGCTCTCAGATCCAGCCCTCTCTCAGCTGCTGCTGCTGCTGCTGCTGCTGCTG 65
21 SerSerProGlyThrGluGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
66 AGCAGCCCAAGACCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCAGCTGAA----- 119
41 IleValValSerGlySerIlePheGlyGlyGlyGlySerLeuValProArgGlySerGly 60
120 ATCAACGAGCTGGTCTGCTAGCGAGGGGGGGGAGC-----GGCGGA 164
61 GlyGlyGlySerGluArgHisPheValValGlnPheLeGlyGluCysTyrThrAen 80
165 GGGGGAATCCGAAAGGCATTTCTGTTCCAGTTCAAGGGGGAGTGTCTACTACACCAAC 224
81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
225 GGGACGCGCGCATACGGCTCGTGACCATATCTACACCGGGAGGAGTACGTGCGC 284

Qy 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db 285 TACGACAGCGAGCTGGCGAGTACCGCGCGTACCGAGCTCGGCGCCAGACGCCGAG 344
Qy 121 TyrTrpAenSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAACACCGCCAGCGAGATCTCGAGAGCAACGCGGGCGAGGTGGACACGGCGTGC 404
Qy 141 ArgHisAenTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAen 160
Db 405 AGACACAACTACGAGGGCGGAGACCAAGACCTCTCCCTCGCGGGCTTGAACAGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAenHisHisAenThrLeuValCysSer 180
Db 465 GTGCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCAACACACTCTGTGTCTGTCG 524
Qy 181 ValThrAspPheTyrProAlaIleValIleValIleValArgTyrPheArgAenGlyGlnGluGlu 200
Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGGCTGCTGTTTCAGGAAATGGCCAGGAGGAG 584
Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAenGlyAspTyrThrPheGlnValLeu 220
Db 585 AAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTCTG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy 241 SerLeuIleSerProIleThrValGluTyrPargAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCTGGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGCGTGGTGGTTCC 770
RESULT 8
AAT86989
ID AAT86989 standard; DNA; 1382 BP.
XX AAT86989;
XX 27-MAR-1998 (first entry)
XX SCE1 single chain gene.
XX Construction; major histocompatibility complex; MHC; fusion complex;
XX SCE1 single chain gene; ss.
XX Synthetic.
XX Key Location/Qualifiers
XX CDS 5..1382
XX /*tag= a
XX WO9728191-A1.
XX 07-AUG-1997.
XX 30-JAN-1997; 97WO-US001617.
XX 31-JAN-1996; 96US-00596387.
XX (DADE-) DADE INT INC.
XX Rhode PR, Jiao J, Burkhardt M, Wong HC;
XX WPI; 1997-402555/37.
XX P-PSDB; AAW29214.
XX Single chain major histocompatibility complex comprising linked alpha and
XX beta chains - useful for suppressing an immune response to an auto:immune
PT

disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes mellitus, etc.

Example 17, Page 140-141; 217pp; English.

The present sequence was used in the construction of major histocompatibility complex (MHC) fusion complexes

Sequence 1382 BP; 320 A; 373 C; 405 G; 284 T; 0 U; 0 Other;

Instrument Scores:

Parent Sequences:					
1. NO.:	1.3e-98	Length:	1382		
2. Ref.:	1145.00	Matches:	227		
3. Percent Similarity:	87.2%	Conservative:	4		
4. Local Similarity:	85.7%	Mismatches:	24		
5. Identity Match:	70.7%	Indels:	10		
	2	Gaps:	3		

0-048-116B-6 (1-306) x AAT86989 (1-1382)

1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
6 ATGGCTCTGCAGATCCCGACGCTCTCTCTCAGCTGCTGTGGTGGTCTGATGGTGTG 65
21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGCGTGTTCAGCTGCTCAGCGTGA----- 119
41 IleValIleSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
120 ATCAACGAAGCTGTGTCTGCTACCGGAGGGGGCGGAAGC-----GGCGGA 164
61 GlyGlyGlySerGluAArgHisPheValValGlnPheLysGlyGluCysTrpTyrThrAsn 80
165 GGGGGAACTCCGAAGGCAATTTCGTGTCTCAGTTCAGGGCGAGTGTCTACTACACCAAC 224
81 GlyThrGlnAArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
225 GGGACGCGCGCATACGGCTCGTGTGACCATCATCTACAAACGGGAGGAGTACGTGGCG 284
101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
285 TACGACAGCGACGTGGCGGAGTACCGCGGGTGACCGAGCTGGGGGGCGGCAGACGCCGAG 344
121 TyrTrpAsnSerGlnProGluIleLeuGluAArgThrArgAlaGluValAspThrAlaCys 140
345 TACTGGAAACAGCCAGCCGAGATCTCTGGAGCGAACCAGCGGCCAGGTGGACACGCCGTGC 404
141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
405 AGACACAACCTACAGGGGGCGGAGACGAGCACTCTCTGGCGGGCTTGAACAGCCCAAT 464
161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer 180
465 GTGCCCATCTCCCTGTCCAGGACAGAGGGCCCTCAACCAACCAACCACTCTGGTCTGTTCG 524
181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
525 GTGACAGATTTCTACCCAGCAAGATCAAGTCCGTGGTTTCAGGAATGGCCAGGAGGAG 584
201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACCTGGACCTTCCAGAGTCTTG 644
221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
645 GTCATGCTGGAGATGACCCCTCATCAGGGAGAGGTCTACACCTGGCCATGTGGAGCATCCC 704
241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerLeuSerAlaArgSerLys 260
705 AGCCTGAAGAGCCCAATCATCTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
261 GlyGlyGlyGlySer 265

Db
756 GCGGTGGTGGTTC 770

RESULT 9
ACA60744

ID ACA60744 standard; DNA; 1382 BP.

ACA60744;

16-JUN-2003 (first entry)

Mouse MHC I-Ad/Ova 323-339 synthetic gene SCE1.

KW MHC; major histocompatibility complex; gene therapy; fusion complex;
 KW peptide-binding groove; T cell modulation; class II MHC; vaccine;
 KW autoimmune disorder; multiple sclerosis; rheumatoid arthritis;
 KW insulin-dependent diabetes mellitus; myasthenia gravis; immunogen;
 KW chronic allergy; mouse; ds: I-Ad; gene.

50
51
52

OS Musc. sp.
OS Synthetic.

AA
PN
US2002198144-A1.XX
PD
26-DEC-2002XX
PF 06-III.-2001. 2001US-00900379-XX
PP 29-III-1994. 94US-00283302

PR 01-FEB-1995; 95US-00382454.
PR 17-JAN-1997. 97US-00376084

XX
/ 1955 1 23 PM TWO

XXI

PI Chavallaz P, Jiao JJ;

DR WPI; 2003-341126/32.

XIX

PT peptide covalently linked to MHC molecule containing peptide-binding groove, used for suppressing immune response in multiple sclerosis, PT allergies.

XX PS Example 17: Fic 29: 126pp: English:

The invention relates to a major histocompatibility complex (MHC) fusion complex (I) comprising an MHC molecule that contains a peptide-binding groove, and a presenting peptide covalently (e.g. an antigenic peptide) linked to the MHC molecule, where (I) is capable of modulating the activity of a T cell. Also included are a DNA construct coding for the MHC molecule, where the MHC molecule is a class II MHC (e.g. mouse I-Ad or I-A_b, or human HLA-DR1 (human leukocyte antigen-DR1)), a multivalent MHC fusion complex comprising two or more linked complexes, identifying a peptide that can modulate the activity of T cells (involving introducing into host cells cloning vectors that each contain the fusion complex DNA, culturing the host cells under conditions suitable for expression of the MHC fusion complex, and selecting host cells that express MHC fusion complex that modulate the activity of T cells), a single recombinant expression vector comprising DNA that codes for the alpha and beta chains of the fusion complex MHC protein, a single recombinant expression vector comprising DNA that codes for a T cell costimulatory factor and the alpha and beta chains of the MHC fusion complex. The DNA constructs can contain heterologous leader peptide sequences and Kozak sequence for efficient expression of the fusion complex. Also included are inducing an immune response in a mammal (including vaccinating a mammal against a targeted disorder, by administering DNA sequence comprising a fusion complex, or DNA sequence coding for a fusion complex which is a single chain fusion molecule) and suppressing an immune response in a mammal by administering to the mammal a DNA sequence comprising an expression vector, encoding a full length MHC molecule that contains a transmembrane domain, and a presenting peptide that is a T cell receptor (TCR) antagonist or partial

Chavaillaz P, Jiao J;
WPI; 1996-129343/13.
P-PSDB; AAR98905.

Major histocompatibility complex fusion complex for modulating T cell activity - used in the treatment of immune disorders, e.g. multiple sclerosis, IDDM and rheumatoid arthritis.

Example 17; Fig 27; 210pp; English.

AAT17586 encodes a murine MHC fusion complex capable of modulating T cell activity encoded by the vector SSC1. The MHC fusion complex comprises at least one MHC molecule containing a peptide-binding groove and a presenting peptide covalently linked to the MHC molecule and opt. a transmembrane domain. DNA encoding a MHC fusion complex may be cloned into a host cell to express the complex. The transformed cells may then be used to identify peptides that modulate, pref. antagonise, T cell activity. DNA encoding a MHC fusion complex or a single chain fusion molecule may be used to vaccinate a mammal against a targeted disorder. The fusion complexes may be used to suppress an immune response in an animal suffering from an immune disorder e.g. multiple sclerosis, insulin-dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or chronic allergies. The complexes may also be used in the treatment of livestock and pets such as cats and dogs. The MHC fusion complexes can be produced such that they contain a single antigenic peptide including one of known structure, additionally a wide range of peptides can be presented for T cell interaction

Sequence 1385 BP; 316 A; 384 C; 398 G; 287 T; 0 U; 0 Other;

ment Scores:

i. No.: 1.3e-98 Length: 1385
e: 1145.00 Matches: 227
ent Similarity: 87.2% Conservative: 4
Local Similarity: 85.7% Mismatches: 24
y Match: 70.7% Indels: 10
2 Gaps: 3

10-048-116B-6 (1-306) x AAT17586 (1-1385)

- 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
- 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGGTGGTGGTGGTGGTGG 65
- 21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
- 66 AGCAGCCCAAGCAGCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCAGCTGAA----- 119
- 41 IleValValSerGlySerTrpaspGlyGlyGlySerLeuValProargGlySerGly 60
- 120 ATCAACGAGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 164
- 61 GlyGlyGlySerGluArgHisPheValValGlnPheGlyGlyGlyCysTrpThrAsn 80
- 165 GGGGGAACCTCGAAGGATTTCTGTTCCAGTTCAAGGGGAGTGTACTACACCAAC 224
- 81 GlyThrGlnArgIleArgLeuValThrArgTrpIleTyrAsnArgGluGluTyrValArg 100
- 225 GCGAGCGCAGCGATACGGCTCGTGACAGATACATCTACACCGGGAGGAGTACGTGGCG 284
- 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProaspAlaGlu 120
- 285 TAGCAGCGGAGCGTGGGCGAGTACCGCGGGTGACCGAGCTCGGGCGGCGAGCGCGAG 344
- 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
- 345 TACTGGACAGCGCGGAGATCTCTGGAGCGNACCGGGCGGCGAGTGTACACGGGCTGC 404
- 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgLeuGluGlnProAsn 160
- 405 AGACACAACTACGAGGGCGCGAGACAGCACCTCTCGCGGGCTTGAACAGGCCCAAT 464

QY	161	ValAlaIleSerLeuSerArgThrGluAlaLeuAenHisHisAenThrLeuValCysSer	180
Db	465	GTGCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACCCACCAACACTCTGGTCTGTTCG	524
QY	181	ValThrAspPheTyrProAlaLysIleValArgTrpPheArgHenglyGlnGluGlu	200
Db	525	GTGACAGATTCTTACCCAGCAAGATCAAAGTGGCTGGTTCAGGAATGGCCAGGAGGAG	584
QY	201	ThrValGlyValSerSerThrGlnLeuIleArgAenGlyAspTrpThrPheGlnValLeu	220
Db	585	ACAGTGGGGGTCTCATCCACAGCTTATTAGGATGGGGACTGGACCTTCCAGGTCCTG	644
QY	221	ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro	240
Db	645	GTGATGTGGAGATGACCCCTCATCAGGAGAGGTCTACACTGCCATGTGGAGCATCCC	704
QY	241	SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys	260
Db	705	AGCCTGAAGAGGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC	755
QY	261	GlyGlyGlySer 265	
Db	756	GGCGGTGGTGGTTC 770	
RESULT 11			
AAT86987			
ID	AAT86987	standard; DNA; 1385 BP.	
AC	AAT86987;		
DT	27-MAR-1998	(first entry)	
DE	SSC1	single chain gene.	
KW	Construction; major histocompatibility complex; MHC; fusion complex;		
KW	SSC1	single chain gene; ss.	
OS	Synthetic.		
FX	Key	Location/Qualifiers	
FT	CDS	6..1385	
FT		/*tag= a	
PN	W09728191-Al.		
PD	07-AUG-1997.		
PF	30-JAN-1997;	97WO-US001617.	
PR	31-JAN-1996;	96US-00596387.	
XX	(DADE-) DADE INT INC.		
PI	Rhode PR, Jiao J, Burkhardt M, Wong HC;		
XX			
DR	WPI: 1997-402555/37.		
DR	P-PSDB; AAW29212.		
XX	Single chain major histocompatibility complex comprising linked alpha and		
PT	beta chains - useful for suppressing an immune response to an auto:immune		
PT	disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes		
PT	mellitus, etc.		
XX			
PS	Example 17; Page 135-137; 217pp; English.		
CC	The present sequence was used in the construction of major		
CC	histocompatibility complex (MHC) fusion complexes		
XX			
SQ	Sequence 1385 BP; 316 A; 383 C; 399 G; 287 T; 0 U; 0 Other;		
Alignment Scores:			
Pred. No.:	1.3e-98	Length:	1385
Score:	1145.00	Matches:	227

Percent Similarity:	87.2%	Conservative:	4
Local Similarity:	85.7%	Mismatches:	2
Y Match:	70.7%	Indels:	1
	2	Gaps:	3

J-048-116B-6 (1-306) x AAT86987 (1-1385)

1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
6 ATGGCTCTGCAGATCCCAAGCCTCTCTCTCAGCTGCTGTGGTGTCTGTATGGTGGT 65
21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
66 AGCAGGCCAAGACCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCACGCTGNA----- 119
41 IleValValSerGlySerTrpAepGlyGlyGlyGlySerLeuValProArgGlySerGly 60
120 ATCAACGAAGCTGGTCTGTCTAGCGAGGGGGCGGAAGC-----GGCGGA 164
61 GlyGlyGlySerGluArgHisPheValValGlnPheIysGlyGluCysTyrTrpThrAen 80
165 GGGGGAACTCCGAAGAAGCATTTCTGGTGTCCAGTTCAAGGGCGAGTCTACTACACCAAC 224
81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
225 GGGACGAGCGCATACGGCTCTGTGACCATATCATCAACCGGGAGGATACGTGGCG 284
101 TyrAspSerAepValGlyGluTyrArgAlaValThrGluLeuGlyArgProAepAlaGlu 120
285 TACGACAGCGACGTGGCGCAGTACCGCGGTGACCGAGCTGGGGCGGCAGACGCCGAG 344
121 TyrTrpAenSerGlnProGluIleLeuGluAArgThrArgAlaGluValAAspThrAlaCys 140
345 TACTTGAACAGCCAGCGGAGATCTTGGAGCGAACCGGGCCGAGGTGGACACGGCGTGC 404
141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAen 160
405 AGACACAACCTACGAGGGCGGAGACCGACCTCTCTCGGGCGGCTTGAAACAGCCCAAT 464
161 ValAlaIleSerLeuSerArgThrGluAlaLeuAenHisHisAsnThrLeuValCysSer 180
465 GTCGCCATCTCCCTGTCTCAGGACAGAGGGCCCTCAACCCACCACTCTGGTGTGTTCG 524
181 ValThrAepPheTyrProAlaLysIleLysValArgTrpPheArgAenGlyGlnGluGlu 200
525 GTGACAGATTTCTACCCAGCCAGATCAAAAGTCGCTGGTTCAGGAAATGGCCAGGAGAG 584
201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAAspTrpThrPheGlnValLeu 220
585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACCTGGACCTTCCAGGTCTTG 644
221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
645 GTCATGCTGGAGATGACCCCTCATACAGGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerLeuAlaArgSerLys 260
705 AGCTGNAGAGCCCATCATCTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
261 GlyGlyGlyGlySer 265
756 GCGGCTGGTGGTTC 770

T 12
1742

ACA60742 standard; DNA; 1385 BP.

ACA60742;

16-JUN-2003 (first entry)

Mouse MHC I-Ad/Ova 323-339 synthetic gene SSC1.

[illegible]

Percent Scores: 1.3e-98 Length: 1385
 i. No.: 1145.00 Matches: 227
 e: 87.2% Conservative: 4
 : Local Similarity: 85.7% Mismatches: 20
 y Match: 70.7% Indels: 10
 Gaps: 3

10-048-116B-6 (1-306) x ACM60742 (1-1385)
 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 6 ATGGCTCTCAGATCCAGCTCCCTCTCAGCTGCTGCTGCTGCTGCTGCTGCTGCTG 65
 21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAAGCTGCTCAGCTGAA----- 119
 41 IleValValSerGlySerTrpAspGlyGlyGlyGlySerLeuValProArgGlySerGly 60
 120 ATCAAGAAAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 164
 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrTyrThrAsn 80
 165 GGGGAAACTCCGAAAGGCAATTTGGTGTCCAGTTCAAGGGCGAGTGTCTACTACACCAAC 224
 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 225 GGGAGCAGCGCATACGGCTCGTACGACAGATACATCTACACCGGGAGGAGTACGTGCGC 284
 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 285 TAGCAGCAGGAGCTGGGGGAGTACCGCGGCGTGACCGAGCTGGGGGCGGCACAGCGCGAG 344
 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 345 TACTGGACAGCAGCGCGGAGATCCTGGAGCGAAGCGGGCGGAGTGGACACGGCGTGC 404
 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
 405 AGACACACACTACGAGGGCGGAGACAGCAGCCTCCTCGCGCGGCTTGAACAGCCCAAT 464
 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
 465 GTGCGCATCTCCCTGCCAGGACAGAGCGCCCTCAACACCAACACTCTGGTCTGTTCCG 524
 181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGluGlu 200
 525 GTGACAGATTTCTACCCAGCCAGATCAAGTGGCTGTTTCAAGGATGCCAGGAGGAG 584
 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrThrPheGlnValLeu 220
 585 ACAGTGGGGTCTCTCCACACAGCTATTAGGAATGGGAGCTGGACCTTCCAGGTCTCTG 644
 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 645 GTATCTGTGAGATGACCTCTATCAGGGAGAGGTCTACACCTGCCCTGTGGAGCATCCC 704
 241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
 261 GlyGlyGlyGlySer 265
 756 GCGGTGGTGGTTC 770

LT 13
 .7587

AAT17587 standard; DNA, 1508 BP.

AAT17587;

26-SEP-1996 (first entry)

XX Vector SCTI-derived single chain gene encoding MHC fusion complex.
 DE MHC; major histocompatibility complex; PCR; polymerase chain reaction;
 XX T cell activity modulator; antagonist; immune disorder; allergy;
 KW multiple sclerosis; insulin-dependent diabetes mellitus;
 KW rheumatoid arthritis; myasthenia gravis; ds.
 XX Synthetic.
 OS
 XX
 FH Key Location/Qualifiers
 CDS 6..1508
 FT /*tag= a
 FT sig_peptide 6..86
 FT /*tag= b
 FT /label= I-Ad beta chain_leader
 FT /note= "murine MHC class II I-Ad gene beta chain leader
 FT sequence"
 FT 87..137
 FT misc_feature /*tag= c
 FT /label= OVA_323-339
 FT /note= "chicken ovalbumin residues 323-339"
 FT 138..167
 FT misc_feature /*tag= d
 FT /note= "10 residue linker peptide"
 FT 168..452
 FT /*tag= e
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 FT /note= "murine MHC class II I-Ad gene beta-1 domain"
 FT 453..734
 FT misc_feature /*tag= f
 FT /label= I-Ad beta2
 FT /note= "murine MHC class II I-Ad gene beta-2 domain"
 FT 735..806
 FT /*tag= g
 FT /note= "24 residue peptide linker"
 FT 807..1067
 FT /*tag= h
 FT /label= I-Ad alpha1
 FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
 FT 1068..1352
 FT misc_feature /*tag= i
 FT /label= I-Ad alpha2
 FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
 FT 1353..1505
 FT /*tag= j
 FT /label= I-Ad alpha-TM
 FT /note= "murine MHC class II I-Ad gene alpha-transmembrane
 FT domain"
 XX
 PN WO9604314-A1.
 XX
 PD 15-FEB-1996.
 XX
 PF 31-JUL-1995; 95WO-US009816.
 XX
 PR 29-JUL-1994; 94US-00283302.
 PR 01-FEB-1995; 95US-00382454.
 XX
 PA (DADE-) DADE INT INC.
 XX
 PI Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
 PI Chavallaz P, Jiao J;
 XX
 DR WPI; 1996-129343/13.
 DR P-PSDB; AAR98906.
 XX
 PT Major histocompatibility complex fusion complex for modulating T cell
 PT activity - used in the treatment of immune disorders, e.g. multiple
 XX sclerosis, IDDM and rheumatoid arthritis.
 PS Example 17; Fig 28; 210pp; English.
 XX

AAT17587 encodes a murine MHC fusion complex capable of modulating T cell activity encoded by the vector SC1. The MHC fusion complex comprises at least one MHC molecule containing a peptide-binding groove and a presenting peptide covalently linked to the MHC molecule and opt. a transmembrane domain. DNA encoding a MHC fusion complex may be cloned into a host cell to express the complex. The transformed cells may then be used to identify peptides that modulate, pref. antagonise, T cell activity. DNA encoding a MHC fusion complex or a single chain fusion molecule may be used to vaccinate a mammal against a targeted disorder. The fusion complexes may be used to suppress an immune response in an animal suffering from an immune disorder e.g. multiple sclerosis, insulin dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or chronic allergies. The complexes may also be used in the treatment of livestock and pets such as cats and dogs. The MHC fusion complexes can be produced such that they contain a single antigenic peptide including one of known structure, additionally a wide range of peptides can be presented for T cell interaction

Sequence 1508 BP; 337 A; 414 C; 440 G; 317 T; 0 U; 0 Other;

ment Scores: 1.46e-98 Length: 1508
 . No.: 1145.00 Matches: 227
 3: nt Similarity: 87.2% Conservative: 4
 Local Similarity: 85.7% Mismatches: 24
 / Match: 70.7% Indels: 10
 2 Gaps: 3

3-048-116B-6 (1-306) x AAT17587 (1-1508)

1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20
 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGCTGTGCTGCTG 65
 21 SerSerProGlyThrGluGlyCysAanSerIleCysPheSerProSerLeuGluHisPro 40
 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTCAGCTGCTCAGCTGAA----- 119
 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 120 ATCAACGAGCTGTGCTGCTAGCGAGGGGGGGGAGC-----GGCGGA 164
 61 GlyGlyGlySerGluArgHisPheValValGlnPheGlyGlyGluCysTrpThrAan 80
 165 GGGGGAACCTCGAAAGGCAATTCGTGTGTCAGATTCAGGGGCGAGTCTACTACACCAAC 224
 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAanArgGluGluTyrValArg 100
 225 GGGACGCGAGCGCATCGCTCGTACCAGATACATCTCAACCGGGAGGAGTACGTGGCG 284
 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 285 TACGACGCGAGCTGGCGAGTACCGCGGTGACCGAGTCCGGCGCCAGACGCCGAG 344
 121 TyrTrpAanSerGlnProGluLeuLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 345 TACTGGAACAGCAGCCGAGATCTGGAGCGAAGCGGGCGAGGTGGACACGCGGTGC 404
 141 ArgHisAanTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAan 160
 405 AGACACAACTACGAGGGGCGGAGACCAACCTCTCCCTGCGGCGCTTGAACAGGCCCAAT 464
 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAanHisAanThrLeuValCysSer 180
 465 GTCCGCTCTCTCTGTCAGGACAGAGGCCCTCAACACCAACCACTCTGCTGTCTG 524
 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAanGlyGlnGlu 200
 525 GTGACAGATTTCTACCCAGCCCAAGATCAAAAGTGGCGCTGTTTCAGGAATGGCCAGGAG 584
 201 ThrValGlyValSerSerThrGlnLeuIleArgAanGlyAspTrpThrPheGlnValLeu 220
 585 ACAGTGGGGGTCTCATCCACACAGCTTTATTAGGAATGGGGAGCTGGACCTTCCAGGTCTCTG 644

Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 Db 645 GTCATGCTGGAGATGAGCCCTCATCAGGAGAGTCTACACCTGCCATGTGGAGCATCCC 704
 Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755

Qy 261 GlyGlyGlyGlySer 265

Db 756 GCGCGTGTGTGTTCC 770

RESULT 14

AAT86988

ID AAT86988 standard; DNA; 1508 BP.

XX AAT86988;

XX 27-MAR-1998 (first entry)

XX SC1 single chain gene.

XX Construction; major histocompatibility complex; MHC; fusion complex;

XX SC1 single chain gene; ss.

XX Synthetic.

XX Key Location/Qualifiers

XX CDS 6..1508

XX WO9728191-A1.

XX 07-AUG-1997.

XX 30-JAN-1997; 97WO-US001617.

XX 31-JAN-1996; 96US-00596387.

XX (DADE-) DADE INT INC.

XX Rhode PR, Jiao J, Burkhardt M, Wong HC;

XX WPI; 1997-402555/37.

XX P-PSDB; AAW29213.

XX Single chain major histocompatibility complex comprising linked alpha and beta chains - useful for suppressing an immune response to an auto-immune disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes mellitus, etc.

XX Example 17; Page 137-139; 217pp; English.

XX The present sequence was used in the construction of major histocompatibility complex (MHC) fusion complexes

XX Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

Alignment Scores: 1.46e-98 Length: 1508
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116B-6 (1-306) x AAT86988 (1-1508)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20

Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGCTGTGCTGCTGCTG 65

21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
 66 AGACGCCAAGACCTTAAAGTATCTCTCAGGCTGTTACGCTGCTCAGCTGAA-----119
 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 120 ATCAACGAGAGCTGGTCTGCTAGCGGAGGGGGGAGC-----GGCGGA 164
 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrThrAsn 80
 165 GGGGGAATCTCCGAAAGGATTTCTGGTCCAGTTCAGGCGGAGGTCTACTACCAAC 224
 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 225 GGGACCGCAGCATCGCTCGTGACCATATCTACACCGGAGGAGTACGTGCC 284
 101 TyrAspSerAspValGlyGlyTrpArgAlaValThrGluLeuGlyArgProHspAlaGlu 120
 285 TACGACAGGAGCTGGGCGAGTACCGCGGTGACCGAGCTGGGGCGGCAGCGCGAG 344
 121 TyrTrpAsnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 345 TACTGGAACAGCGCGAGATCTCTGAGCGAAGCGGGCGAGGTGACACCGCGTGC 404
 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgGluGlnProAsn 160
 405 AGACACAATAGAGGGGCGGAGACGACCTCTCCGCGGCTTGAACAGCCCAAT 464
 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisGlyAsnThrLeuValCysSer 180
 465 GTCCGATCTCTCTCCAGGACAGAGGCGCTCAACACCAACACACTCTGGTCTGTTG 524
 181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGlu 200
 525 GTGACAGATTTCTACCGACCAAGATCAAGTCCGCTGCTCAGGAATGCCAGGAGGAG 584
 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrThrPheGlnValLeu 220
 585 ACAGTGGGGGTCTCATCCACAGACTTATAGGAATGGGAGCTGGACCTTCCAGGTCTG 644
 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 645 GTCATGCTGGAGATGACCTCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCC 704
 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 705 AGCCTGAAGAGCCCATCATCTGTGGAGTGG-----ACTAGTGGTGGCGTGGCAGC 755
 261 GlyGlyGlyGlySer 265
 756 GGGGGTGGTGGTTC 770

JLT 15

:9069

AAX89069 standard; DNA; 1508 BP.

AAX89069;

14-SEP-1999 (first entry)

Single chain IAD/OVA 323-229 MHC fusion protein encoding DNA.

Major histocompatibility complex; MHC; single chain MHC; sc-MHC; Ig; peptide binding groove; immunoglobulin; T cell receptor; immune response; immune-related disorder; antigenic peptide; fusion protein; ss.

Synthetic.

W09921572-A1.

06-MAY-1999.

13-OCT-1998; 98WO-US021520.

XX 29-OCT-1997; 97US-00960190.
 PR (SUNO-) SUNOL MOLECULAR CORP.
 XX Rhode PR, Acevedo J, Burkhardt M, Jiao J, Wong HC;
 XX WPI; 1999-418411/35.
 DR P-PSDB; AAY27111.
 DR Single chain major histocompatibility complex class I complexes.
 XX Example 1; Fig 1; 148pp; English.
 XX The invention relates to new single chain major histocompatibility
 CC complex (sc-MHC) class II complexes that comprise a peptide binding
 CC groove, and a modified class II beta 2 chain or covalently linked
 CC immunoglobulin (Ig) light chain constant (CI) region. The MHC complexes
 CC are useful for detection and analysis of peptide ligands, pathogenic T-
 CC cells, for functional, cellular and molecular assays. They can be used to
 CC identify and isolate T cell receptor and/or MHC agonists and antagonists.
 CC They can be used in vivo to compete with pathogenic antigen presenting
 CC cells involved in immune-related disorders. They can also be used to
 CC raise antibodies and to screen immune cells. It is also use in a method
 CC of suppressing an immune response in mammals. The sc-MHC complexes
 CC comprising modified class II beta 2 chains and/or Ig-CI regions are
 CC soluble and provide enhanced yield. These MHC complexes also can contain
 CC single antigenic peptides readily isolated from expressing cells in
 CC significant quantities. The polyspecific MHC complexes also provide a
 CC means to detect cells expressing multiple target structures with a single
 CC complex. The present sequence represents a DNA encoding a single chain
 CC IAD/OVA 323-229 MHC fusion protein
 XX

SQ Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	1,46e-98	Length:	1508
Score:	1145.00	Matches:	227
Percent Similarity:	87.2%	Conservative:	4
Best Local Similarity:	85.7%	Mismatches:	24
Query Match:	70.7%	Indels:	10
DB:	2	Gaps:	3

US-10-048-116B-6 (1-306) x AAX89069 (1-1508)

QY 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db 6 ATGGCTGTGACATCCAGCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTGTG 65
 QY 21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGACGCCAAGACCTTAAAGTATCTCTCAGGCTGTTACGCTGCTCAGCTGAA-----119
 QY 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db 120 ATCAACGAGAGCTGGTCTGCTAGCGGAGGGGGGAGC-----GGCGGA 164
 QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrThrAsn 80
 Db 165 GGGGGAATCTCCGAAAGGATTTCTGGTCCAGTTCAGGCGGAGGTCTACTACCAAC 224
 QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 Db 225 GGGACCGCAGCATCGCTCGTGACCATATCTACACCGGAGGAGTACGTGCC 284
 QY 101 TyrAspSerAspValGlyGlyTrpArgAlaValThrGluLeuGlyArgProHspAlaGlu 120
 Db 285 TACGACAGGAGCTGGGCGAGTACCGCGGTGACCGAGCTGGGGCGGCAGCGCGAG 344
 QY 121 TyrTrpAsnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 Db 345 TACTGGAACAGCGCGAGATCTCTGAGCGAAGCGGGCGAGGTGACACCGCGTGC 404

```

141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
|||||
405 AGACACAACACTAGGAGGGCGGAGACACACCTCCCTGGGGGGCTTGAACAGCCCAAT 464
|||||
161 VallalalleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer 180
|||||
465 GTCGCCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCACCAACACACTCTGGTCTGTTCG 524
|||||
181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGlu 200
|||||
525 GTGACAGATTCTACCCAGCCCAAGATCAAGTGCCTGCTTCAGGAATGGCCAGGAGAG 584
|||||
201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrThrPheGlnValLeu 220
|||||
585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCCTG 644
|||||
221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
|||||
645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTTTACACCTGCTGATGTGGAGCATCCC 704
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241 SerLeuLysSerProIleThrValGluTyrPargAlaGlnSerGluSerAlaArgSerLys 260
|||||
705 AGCCTGAAGAGCCCCCATCACTGTGGAGTGG-----ACTAGTGTGGGGGTGGCAGC 755
|||||
261 GlyGlyGlyGlySer 265
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756 GCGGTGGTGGTTCC 770
|||||

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 Maximum Match 100%
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 3RN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
 3APOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

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 3: gb_ph.*
 4: gb_pl.*
 5: gb_pr.*
 6: gb_ro.*
 7: gb_ats.*
 8: gb_sy.*
 9: gb_un.*
 10: gb_vi.*
 11: gb_ov.*
 12: gb_hcg.*
 13: gb_in.*
 14: gb_om.*
 15: gb_ba.*

Pred. No. is the number of results predicted by chance to have a
 score greater than or equal to the score of the result being printed,
 and is derived by analysis of the total score distribution.

SUMMARIES

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1	1496	100.0	1484	2 AX081280
2	1182	79.0	978	6 MUSHHIAA
3	1182	79.0	4713	2 AR199665

4	1161	77.6	771	6	AY452201	Mus muscu
5	1107	74.0	1085	6	BC029620	Mus muscu
6	1100	73.5	1110	6	BC043925	Mus muscu
7	1078	72.1	776	2	AR106256	Sequence
8	1078	72.1	776	2	AR229608	Sequence
9	1078	72.1	776	2	AR363023	Sequence
10	1078	72.1	1084	6	BC019721	Mus muscu
11	1078	72.1	1109	6	BC031711	Mus muscu
12	1075	71.9	978	6	MMMH01	Mouse fragm
13	1075	71.9	978	6	MUSHHIAA	M21931 Mouse MHC c
14	1069	71.5	776	2	AR365183	Sequence
15	1065.5	71.2	1508	2	AR033963	Sequence
16	1065.5	71.2	1508	2	AR152030	Sequence
17	1065.5	71.2	1508	2	AR175096	Sequence
18	1065.5	71.2	1508	2	CS079300	Sequence
19	1065.5	71.2	1508	2	AX032544	Sequence
20	1057	70.7	942	6	MMTEANON	X52643 Mouse mRNA
21	1056.5	70.6	1508	2	BD138632	Soluble M
22	1048	70.1	932	6	MUSHHIAA	K01922 Mouse MHC c
23	1048	70.1	942	6	AF119253	Mus muscu
24	1048	70.1	942	6	AF119254	Mus muscu
25	1020	68.2	998	6	MUSHHIAA	K01924 Mouse MHC c
26	1007.5	67.3	1382	2	AR033964	Sequence
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31	996	66.6	828	6	AF065910	Mus muscu
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33	993.5	66.4	1385	2	MUSHHIAA	M11358 Mouse MHC c
34	993.5	66.4	1385	2	AR033962	Sequence
35	993.5	66.4	1385	2	AR175095	Sequence
36	993.5	66.4	1385	2	CS079299	Sequence
37	993.5	66.4	1385	2	AX032543	Sequence
38	993	65.7	1243	2	AX490802	Sequence
39	976	65.2	886	6	MUSHHIAA	M11357 Mouse MHC c
40	966	64.6	771	6	AY626198	Rattus no
41	966	64.6	771	6	AY626199	Rattus no
42	964	64.4	771	6	AF307302	Rattus no
43	964	64.4	771	6	AY626195	Rattus no
44	964	64.4	771	6	AY626196	Rattus no
45	964	64.4	795	6	AY701537	Rattus no

ALIGNMENTS

RESULT 1	AX081280	AX081280	Sequence 1 from Patent WO0109194.	1484 bp	DNA	linear	PAT 27-FEB-2001
LOCUS	AX081280	Sequence 1 from Patent WO0109194.					
DEFINITION	AX081280	Sequence 1 from Patent WO0109194.					
ACCESSION	AX081280	Sequence 1 from Patent WO0109194.					
VERSION	AX081280.1	GI:13170129					
KEYWORDS							
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ORGANISM							
REFERENCE							
AUTHORS							
TITLE							
JOURNAL							
FEATURES							
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CDS							

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CPANLGLGSPVFIFFPKIKQVLMISLSPITVTVVVDSEDDPDVQISFVNNVEVHT
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CIN

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ie:	1496.00	Matches:	278
cent Similarity:	100.0%	Conservatives:	0
t Local Similarity:	100.0%	Mismatches:	0
y Match:	100.0%	Indels:	0
	2	Gaps:	0
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1	ATGCCGTGCAGCAGAGCTCTGATTCTGGGGGTCTCTGCCCTGAACACCATGCTCAGCCTC	60	
21	CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyrGlyThrThrValTyr	40	
61	TCCGAGGTTGAAGACACATTGAGCCGACACAGTAGGCTTCTATGGTACACCTGTTAT	120	
41	GlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr	60	
121	CAGTCTCCTGGAGACATTGGCCAGTACACACATGAATTGATGGTGGTGGTCTCTAT	180	
61	ValAspLeuAspIleGlyThrValTyrArgLeuProGluPheGlyGlnLeuLeuLeu	80	
181	GTGGACTTGGATGAAGAAGAAACTGTCTGGAGGCTCTCTGGAGTTGGCCAAATTGATCTC	240	
81	PheGluProGlnGlyGlyLeuGlnAsnIleAlaGluLysHisAsnLeuGlyIleLeu	100	
241	TTTGAGCCCCAAGGTGGACTGCAGAAACATAGCTGCAGAAAACACACTGGGAATCTTG	300	
101	ThrLysArgSerAsnPheThrProAlaThrAsnGluAlaProGlnAlaThrValPhePro	120	
301	ACTAAGAGGTCAAATTTACCCCGAGTACCAATGAGGCTCTCAAGCGAGCTGTGTCCCC	360	
121	LysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePhe	140	
361	AAGTCCCCCTGTCTGCTGGGTGAGCCCAACACCTTATCTGTTGTGGACAACTCTTC	420	
141	ProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyr	160	
421	CCACCTGTGATCAACATCATGCTGCAGAAATAGCAAGTCACTCAGACGCGCTTAT	480	
161	GluThrSerPheLeuValAsnArgAspHisSerPheHisLysLeuSerTyrLeuThrPhe	180	
481	GAGACGAGCTTCTCTGCTCAACCGTGACCATCTCTTCCACAGCTGTCTTATCTCACTTC	540	
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541	ATCCCTTCTGTGATGACATTTATGACTGCAAGGTGAGCACTGGGGCTTGAGGAGCGG	600	
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221	GlyGlyGlySerThrThrAlaProSerAlaGlnLeuGluLysGluLeuGlnAlaLeuGlu	240	
661	GGTGGAGGATCCACTACAGCTCCATCAGCTCAGTCCGAAAAAGAGCTCCAGGCCCTGGAG	720	
241	LysGluAsnAlaGlnLeuGluTrpGluLeuGlnAlaLeuGluLysLeuLeuAlaGlnAla	260	
721	AAGGAAATGACAGCTGGATGGGAGTTGCAAGCACTGGAAAGGAACCTGGCTCAGGCA	780	

Qy	261	AlaSerGluProArgGlyProThrIleLysProCysProProCysLysCysPro 278	
Db	781	GCATCTGAGCCAGAGGGCCCAATCAATCAAGCCCTGTCTCCATGCAATGCCCA 834	
RESULT 2			
MUSMHAAD			
LOCUS			
DEFINITION	MUSMHAAD	978 bp	mRNA linear ROD 27-APR-1993
LOCUS	Mouse MHC class II H2-IA-alpha gene (d haplotype) mRNA, complete cds		
ACCESSION	K01923		
VERSION	K01923.1	GI:199449	
KEYWORDS	antigen; cell surface glycoprotein; class II gene; glycoprotein; histocompatibility antigen; integral membrane protein; major histocompatibility complex.		
SOURCE	Mus musculus (house mouse)		
ORGANISM	Mus musculus		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi; Muroidea; Muridae; Murinae; Mus.		
REFERENCE	1	(bases 1 to 978)	
AUTHORS	Benoist,C.O., Mathis,D.J., Kanter,M.R., Williams,V.E. II and McDevitt,H.O.		
TITLE	Regions of allelic hypervariability in the murine A alpha immune response gene		
JOURNAL	Cell 34 (1), 169-177 (1983)		
PUBMED	6309407		
COMMENT	Original source text: Mus musculus (strain BALB/c, sub_species domesticus) spleen cDNA to mRNA. The protein domains are as follows: first external protein domain (D1) at bases 93-356; second external protein domain (D2) at bases 357-638; connecting peptide, transmembrane region, and cytoplasmic tail (CP, TM, C) at bases 639-791. [1] also sequenced the IA-alpha genes from mice of b, f, u and q haplotypes.		
FEATURES	Location/Qualifiers		
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	/product="IA-alpha mRNA"		
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	/gene="MHC H2-IA-alpha"		
	/product="unnamed"		
ORIGIN	44 bp upstream of HinfI site, chromosome 17.		
Alignment Scores:			
Pred. No.:	2,12e-114	Length:	978
Score:	1182.00	Matches:	219
Percent Similarity:	100.0%	Conservative:	0
Best Local Similarity:	100.0%	Mismatches:	0
Query Match:	79.0%	Indels:	0
DB:	6	Gaps:	0

4-2-11

-10-048-116B-2_COPY_1_278 (1-278) x MUSMHIAD (1-978)

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41 GlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeuPheTyr 60
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61 ValAspLeuAspPlyLysLysThrValTyrArgLeuProGluPheGlyGlnLeuLeu 80
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81 PheGluProGlnGlyLeuGlnAsnIleAlaAlaGluLysHisAsnLeuGlyIleLeu 100
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L99665
US
FINITION
SSION
SION
WORDS
ORCE
ORGANISM
Unclassified.
REFERENCE
1 (bases 1 to 4713)
Webb,S.R., Wingvist,O., Karlsson,L., Jackson,M.R. and Peterson,P.A.
MHC class II antigen-presenting systems and methods for activating
CD4+ T cells
Patent: US 6355479-A 7 12-MAR-2002;
Location/Qualifiers
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y Match: 79.0% Indels: 0
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21 CysGlyGlyGluAspAspIleGluAlaAspHisValGlyPheTyGlyThrThrValTyr 40
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121 LysSerProValLeuLeuGlyGlnProAsnThrLeuIleCysPheValAspAsnIlePhe 140
819 AAGTCCCTGTCTGCTGGGTGAGCCCAACACCTTATCTGCTTTGTGGACAACATCTTC 878
141 ProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyr 160
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999 ATCCCTTCTGATGATGACATTTATGACTGCAAGGTGGAGCACTGGGGCCTGGAGGAGCG 1058
201 ValLeuLysHisTrpGluProGluIleProAlaProMetSerGluLeuThrGluThr 219
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LOCUS Mus musculus H-2 class II histocompatibility antigen, A-D alpha
DEFINITION chain precursor, mRNA, complete cds.
ACCESSION AY452201
VERSION AY452201.1 GI:38373608
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 771)
AUTHORS Gao,M., Wang,H. and Wang,Q.
TITLE Establishment of pIRES- I-A(d) alpha beta and stable expression of
BALB/c mouse I-Ad alpha beta chain gene in NIH3T3 cell line
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 771)
AUTHORS Gao,M., Wang,H. and Wang,Q.
TITLE Direct Submission
JOURNAL Submitted (29-OCT-2003) Lab of Transfusion Transferred virus,
Transfusion Institute Beijing, 27 Taiping Road, Beijing 100039, PR
China
FEATURES Location/Qualifiers
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RESULT 5

BC029620

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

PUBMED

REFERENCE

AUTHORS

TITLE

JOURNAL

REMARK

COMMENT

FEATURES

source

BC029620 1085 bp mRNA linear ROD 04-OCT-2003
Mus musculus histocompatibility 2, class II antigen A, alpha, mRNA
(CDNA clone MGC:25392 IMAGE:2609494), complete cds.

BC029620

BC029620.1 GI:20987326

MGC.

Mus musculus (house mouse)

Mus musculus

Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 1085)

Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
Klausner, R.D., Collins, P.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
Scheetz, T.E., Brownstein, M.J., Ustin, T.B., Toshiyuki, S.,
Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,
Aramson, R.D., Mullany, S.J., Bosak, S.A., McEwan, P.J.,
McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,
Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,
Villalon, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,
Fahey, J., Helton, E., Kettman, M., Madan, A., Rodriguez, S.,
Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y.,
Bouffard, G.G., Blakeley, R.W., Touchman, J.W., Green, E.D.,
Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M.,
Butterfield, Y.S., Krzywinski, M.I., Skalek, U., Smalls, D.E.,
Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.

Generation and initial analysis of more than 15,000 full-length

human and mouse cDNA sequences

Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)

12477932

2 (bases 1 to 1085)

Strausberg, R.

Direct Submission

Submitted (06-MAY-2002) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,

USA

NIH-MGC Project URL: <http://mgc.nci.nih.gov>

Contact: MGC help desk

Email: cgaps-remail.nih.gov

Tissue Procurement: Jeffrey Green M.D.

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Baylor College of Medicine Human Genome

Sequencing Center

Center code: BCM-HGSC

Web site: <http://www.hgsc.bcm.tmc.edu/cdna/>

Contact: amgobcm.tmc.edu

Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Loulsegod, H.,

Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati,

A.N., Gibbs, R.A.

Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>

Series: IRAP Plate: 30 Row: a Column: 2

This clone was selected for full length sequencing because it

passed the following selection criteria: matched mRNA gi: 13540710.

Location/Qualifiers

1..1085

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Alignment Scores:
1. No.: 1.94e-106 Length: 1085
2. re: 1107.00 Matches: 205
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4. st Local Similarity: 93.6% Mismatches: 11
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6. Gaps: 0

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271 TTTGACCCCAAGTGGAGCTACAAACATAGCTACAGGAAACACAACTTGGAGGCTGG 330
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391 AAGTCCCTGTGCTGTGGGTGACGCCCAATACCTTATCTGCTTTGTGGACAACTCTTC 450
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141 ProProValIleAsnIleThrTrpLeuArgAsnSerLysSerValThrAspGlyValTyr 160
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RESULT 6
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Mus musculus histocompatibility 2, class II antigen A, alpha, mRNA
(cDNA clone MGC:49437 IMAGE:4023996), complete cds.
BC043925
BC043925.1 GI:27882597
MGC.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridea; Muridae; Murinae; Mus.
1 (bases 1 to 1110)
Straussberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Heideh,F.,
Diatchenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Stapleton,M., Soares,M.B., Bonaldo,M.P., Casavant,T.L.,
Schetz,T.E., Brownstein,M.J., Usdin,T.B., Toshlyuki,S.,
Carninci,P., Prange,C., Raha,S.S., Loquellano,N.A., Peters,G.J.,
Abramson,R.D., Mullahy,S.J., Bosak,S.A., McEwan,P.J.,
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Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
Villalon,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
Fahey,J., Helton,E., Kettman,M., Madan,A., Rodriguez,S.,
Sanchez,A., Whiting,M., Madan,A., Young,A.C., Shevchenko,Y.,
Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smallus,D.E.,
Schmerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Mammalian Gene Collection Program Team
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
12477932
2 (bases 1 to 1110)
NIH MGC Project
Direct Submission
Submitted (10-JAN-2003) National Institutes of Health, Mammalian
Gene Collection (MGC), Bethesda, MD 20892-2590, USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgapbs@mail.nih.gov
Tissue Procurement: Lohar Hennighausen Ph.D., Robin Humphreys
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc.mgc@nih.gov
Akhter,N., Avdey,K., Beckstrom-Sternberg,S.M., Benjamin,B.,
Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C., Brooks,S.,
Dietrich,N.L., Granite,S., Guan,X., Gupta,J., Haghighi,P.,
Hansen,N., Ho,S.-L., Karlins,E., Kwong,P., Latic,P., Legaspi,R.,
Maduro,Q.L., Mastello,C., Maskeri,B., Mastrian,S.D., McCloskey,J.C.,
McDowell,J., Pearson,R., Stantropop,S., Thomas,P.J., Touchman,J.W.,

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Tsurgeon, C., Vogt, J.L., Walker, M.A., Wetherby, K.D., Wiggins, L., Young, A., Zhang, L.-H. and Green, E.D.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/ILN at: <http://image.llnl.gov>
Series: IRAP Plate: 85 Row: f Column: 12
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 31981715.

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gene

CDS

RESULT 7
LOCUS ARI06256 776 bp DNA linear PAT 14-FEB-2001
DEFINITION Sequence 4 from patent US 6106840.
ACCESSION ARI06256
VERSION ARI06256.1 GI:12820786
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 776)
AUTHORS Clark, B.R., Sharma, S.D. and Lerch, B.L.
TITLE MHC conjugates useful in ameliorating autoimmunity
JOURNAL Patent: US 6106840-A 4 22-AUG-2000;
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/organism="unknown"
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Alignment Scores:
Pred. No.: 1.36e-103 Length: 776
Score: 1078.00 Matches: 201
Percent Similarity: 94.1% Conservative: 5
Best Local Similarity: 91.8% Mismatches: 13
Query Match: 72.1% Indels: 0
Gaps: 0
DB:
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546 ATCCCTTCTGACGATGACATTTATGACTGCAAGGTGGAAACACTGGGGCCTGGAGGAGCG 605
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BULT 8

229608 AR229608 776 bp DNA linear PAT 20-DEC-2002

DEFINITION Sequence 4 from patent US 6451314.

ACCESSION AR229608

VERSION AR229608.1 GI:27269264

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 776)

AUTHORS Clark, B.R., Sharma, S.D. and Lerch, B.L.

TITLE MHC conjugates useful in ameliorating autoimmunity

JOURNAL Patent: US 6451314-A 4 17-SEP-2002;

Anergen, Inc.; Seattle, WA

FEATURES

Location/Qualifiers

1..776

/organism="unknown"

/mol_type="genomic DNA"

ORIGIN

Alignment Scores:
Pred. No.: 1.36e-103 Length: 776
Score: 1078.00 Matches: 201
Percent Similarity: 94.1% Conservative: 5
Best Local Similarity: 91.8% Mismatches: 13
Query Match: 72.1% Indels: 0
DB: 2 Gaps: 0

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DEFINITION Sequence 5 from patent US 5194425.

ACCESSION AR363023

VERSION AR363023.1 GI:34423771

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 776)

AUTHORS Sharma, S.D., Lerch, L.B. and Clark, B.R.

TITLE MHC-mediated toxic conjugates useful in ameliorating autoimmunity

JOURNAL Patent: US 5194425-A 5 16-MAR-1993;

Anergen, Inc.; Redwood City, CA

FEATURES

Location/Qualifiers

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/organism="unknown"

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ORIGIN

Alignment Scores:
Pred. No.: 1.36e-103 Length: 776
Score: 1078.00 Matches: 201
Percent Similarity: 94.1% Conservative: 5
Best Local Similarity: 91.8% Mismatches: 13
Query Match: 72.1% Indels: 0
DB: 2 Gaps: 0

US-10-048-116B-2_COPY_1_278 (1-278) x AR363023 (1-776)

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Db 66 TGTGGAGGTGAACGACATTTGAGCCGACACGCTAGGCACCTATGATATAGTGATAT 125
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* LT 10
9721
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NITION
iSSION
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iGRDS
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iGANISM
iRENCE
iTHORS

BC019721 1084 bp mRNA linear ROD 30-JUN-2004
Mus musculus histocompatibility 2, class II antigen A, alpha, mRNA
(CDNA clone MGC:30249 IMAGE:3669693), complete cds.
BC019721
BC019721.1 GI:18043825
MGC.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridea; Muridae; Murinae; Mus.
1 (bases 1 to 1084)
Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,F.,
Diachenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Stapleton,M., Soares,M.B., Bonaldo,M.P., Casavant,T.L.,
Scheetz,T.E., Brownstein,M.J., Ustin,T.B., Toshiyuki,S.,
Carninci,P., Frange,C., Raha,S.S., Loquellano,N.A., Peters,G.J.,
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Villalon,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
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Butterfield,Y.S., Krzyzanski,M.I., Skalska,U., Smailus,D.E.,
Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
12477932
2 (bases 1 to 1084)
Strausberg,R.
Direct Submission
Submitted (19-DEC-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Lothar Hennighausen Ph.D., Robin Humphreys
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CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Baylor College of Medicine Human Genome
Sequencing Center
Center code: BCM-HGSC
Web site: <http://www.hgsc.bcm.tmc.edu/cdna/>
Contact: amgobcm.tmc.edu
Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Loulaeeged, H.,
Kovis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati,
A.N., Gibbs, R.A.

Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Series: IRAK Plate: 40 Row: d Column: 4
This clone was selected for full length sequencing because it
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FEATURES

Location/Qualifiers
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ORIGIN

Alignment Scores:
Pred. No.: 2,2e-103 Length: 1084
Score: 1078.00 Matches: 201
Percent Similarity: 94.1% Conservative: 5
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486 GAGACCAAGCTTCTCGTCAACCGTGACTATTCTTCCACAAGCTGCTTATCTCACCITC 545
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546 ATCCCTTCTGACGATGACATTTATGACTGCAAGGTGGAACACTGGGGCCTGGAGGACGG 605
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3963
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NTION Sequence 122 from patent US 5869270.
SSION AR033963
ION AR033963.1 GI:5949568
* ORDS
CE Unknown.
GANISM Unclassified.
* RENGE 1 (bases 1 to 1508)
* THORS Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
* TLE Single chain MHC complexes and uses thereof
* URNAL Patent: US 5869270-A 122 09-FEB-1999;
* URES Location/Qualifiers
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* t Local Similarity: 98.5% Mismatches: 0
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792 GCGGGTTCCTCGAGTGAACGACGACATTGAGCGCGACCGTAGGCTTCTATGTTACAACT 851
39 ValTyrGlnSerProGlyAspIleGlyGlnTyrThrHisGluPheAspGlyAspGluLeu 58
852 GTTTATCAGTCTCCTGAGACATTTGGCCAGTACACATGTAATTTGATGGTATGAGTTG 911
59 PheTyrValAspLeuAspLysLysValThrValTyrArgLeuProGluPheGlyGlnLeu 78
912 TTCTATGTGAGCTTGGATGAAGAGAAACTGTCTGGAGGCTTCCCTGAGTTTGGCCAAATTG 971
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GenCore version 5.1.8
Copyright (c) 1993 - 2006 Bioceleration Ltd.

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Maximum Match 100%

Listing first 45 summaries

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- 14: gb_om.*
- 15: gb_ba.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	1154	71.2	1013	2	AR047957	Sequence
5	1145	70.7	1382	2	AR033964	Sequence
6	1145	70.7	1382	2	AR175097	Sequence
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29	1041.5	64.3	792	6	MUSMHIAB5	M13540 Mouse MHC c
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31	1036.5	64.0	792	6	AF119252	Mus muscu
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33	1030.5	63.6	1078	6	AF293061	Mus muscu
34	1028.5	63.5	792	6	MUSMHIABK	M13538 Mouse MHC c
35	1016.5	62.7	792	6	MUSMHIABU	M13539 Mouse MHC c
36	998.5	61.6	1070	6	AF015280	Mus muscu
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ACCESSION	AX081281	AX081281	Sequence 2 from Patent WO0109194.	921 bp	DNA	linear	PAT 27-FEB-2001
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Glaiichenhaus, N. and Malherbe, L.
Recombinant proteins and molecular complexes derived therefrom,
analogous to molecules involved in immune responses
Patent: WO 0109194-A 2 08-FEB-2001;
CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (PR)

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721 AGCCTGAAGAGGCCCATCACTGTGTGAGTGGAGGGCACAGTCCGAGTCTGCGCGGAGCAAG 780  
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ACCESSION AR047947  
VERSION AR047947.1 GI:5970290  
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ORGANISM Unknown.  
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AUTHORS Kappler,J.W. and Marrack,P.  
TITLE Product and process for T cell regulation  
JOURNAL Patent: US 5820866-A 25 13-OCT-1998;  
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Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160  
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LT 5
3964
S
NITION
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Unknown.
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Single chain MHC complexes and uses thereof
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Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGGCTGTTTCAGGAATGGCCAGGAGGAG 584

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RESULT 6

ARI75097

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

ARI75097 1382 bp DNA linear PAT 17-DEC-2001

Sequence 123 from patent US 6309645.

ARI75097

ARI75097.1 GI:17916396

Unknown.

Unknown.

Unclassified.

1 (bases 1 to 1382)

Rhode, P.R., Jiao, J.-A., Burkhardt, M. and Wong, H.C.

MHC molecules and uses thereof

Patent: US 6309645-A 123 30-OCT-2001;


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               Matches: 227
               Conservative: 4
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               Indels: 10
               Gaps: 3
               Percent Similarity: 87.2%
               Best Local Similarity: 85.7%
               Query Match: 70.7%
               DB: 2
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21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
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41 IleValValSerGlySerTrpAaspGlyGlyGlySerLeuValProArgGlySerGly 60
120 ATCAACGAGACTGCTGCTGCTAGCGAGGGGGCGGAAGC-----GGCGGA 164
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DEFINITION    Sequence 123 from Patent EP1526141.
ACCESSION     CS079301
VERSION       CS079301.1      GI:63093743
KEYWORDS      unidentified
SOURCE        unidentified
ORGANISM      unclassified sequences
REFERENCE     1
AUTHORS       Rhode, P., Jiao, J.A., Burkhardt, M. and Wong, H.C.
TITLE         MHC complexes and uses thereof
JOURNAL       Patent: EP 1526141-A 123 27-APR-2005;
              Altor Bioscience Corporation (US)
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Score:          1145.00
Percent Similarity: 87.2%
Best Local Similarity: 85.7%
Query Match:    70.7%
DB:             2
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QY 21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
DB 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAAGCTGCTCAGCTGAA----- 119
QY 41 IleValValSerGlySerTrpAaspGlyGlyGlySerLeuValProArgGlySerGly 60
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AX032545 1382 bp DNA linear PAT 20-SEP-2000
 Sequence 123 from Patent EP0997477.
 AX032545
 AX032545.1 GI:10279486
 unclassified
 unclassified sequences.
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 CHAVAILLAZ,P.A., EDWARDS,A.C., GRAMMER,S., JIAO,J.A., RHODE,P.R.,
 WEIDANZ,J.A. and WONG,H.C.
 Mhc complexes and uses thereof
 Patent: EP 0997477-A 123 03-MAY-2000;
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JRES
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 . NO.: 1.91e-110 Length: 1382
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 Local Similarity: 85.7% Mismatches: 24
 y Match: 70.7% Indels: 10
 2 Gaps: 3

J-048-116B-6 (1-306) x AX032545 (1-1382)

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RESULT 9
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 LOCUS
 DEFINITION Sequence 121 from patent US 5869270.
 ACCESSION AR033962
 VERSION AR033962.1 GI:5949567
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 1385)
 AUTHORS Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
 TITLE Single chain MHC complexes and uses thereof
 JOURNAL Patent: US 5869270-A 121 09-FEB-1999;
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 Location/Qualifiers
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 /organism="unknown"
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ORIGIN

Alignment Scores:
 Pred. No.: 1.91e-110 Length: 1385
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116B-6 (1-306) x AR033962 (1-1385)

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AR175095 1385 bp DNA linear PAT 17-DEC-2001
SEQUENCE 121 from patent US 6309645.

AR175095.1 GI:17916394
Unknown.
Unclassified.
REFERENCE 1 (bases 1 to 1385)
AUTHORS Rhode, P.R., Jiao, J.-A., Burkhardt, M. and Wong, H.C.
TITLE MHC molecules and uses thereof
JOURNAL Patent: US 6309645-A 121 30-OCT-2001;
FEATURES Location/Qualifiers
source 1..1385
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Alignment Scores:
i. No.: 1..91e-110 Length: 1385
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: Local Similarity: 85.7% Mismatches: 24
y Match: 70.7% Indels: 10
2 Gaps: 3
10-048-116B-6 (1-306) x AR175095 (1-1385)

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DEFINITION Sequence 121 from Patent EPI526141.
ACCESSION CS079299
VERSION CS079299.1 GI:63093741

KEYWORDS
SOURCE unidentified
ORGANISM unidentified
REFERENCE 1
AUTHORS Rhode, P.R., Jiao, J.-A., Burkhardt, M. and Wong, H.C.
TITLE MHC complexes and uses thereof
JOURNAL Patent: EP 1526141-A 121 27-APR-2005;
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Local Similarity: 85.7% Mismatches: 24
/ Match: 70.7% Indels: 10
2 Gaps: 3

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66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTACAGCTGCTCAGCTGAA----- 119
41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
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61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrTrpAsn 80
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LOCUS AX032543 1385 bp DNA linear PAT 20-SEP-2000
DEFINITION Sequence 121 from Patent EP0997477.
ACCESSION AX032543
VERSION AX032543.1 GI:10279484
KEYWORDS
SOURCE unidentified
ORGANISM unclassified sequences.
REFERENCE
AUTHORS Chavallaz, P.A., Edwards, A.C., Grammer, S., Jiao, J.A., Rhode, P.R.,
Weidanz, J.A. and Wong, H.C.
Mhc complexes and uses thereof
Patent: EP 0997477-A 121 03-MAY-2000;
SUNOL MOLECULAR CORP (US)
JOURNAL
FEATURES
Location/Qualifiers
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/organism="unidentified"
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ORIGIN
Alignment Scores: 1.91e-110 Length: 1385
Pred. No.: 1145.00 Matches: 227
Score: 87.2% Conservat: 4
Percent Similarity: 85.7% Mismatches: 24
Best Local Similarity: 70.7% Indels: 10
Query Match: 2 Gaps: 3
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US-10-048-116B-6 (1-306) x AX032543 (1-1385)

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Qy 101 TyrAspSerAspValGlyGlyArgHisPheValValGlnPheLysGlyGlyCysTyrTrpAsn 120
Db 285 TACGACAGGACGCTGGGCGAGTACCGCGGTGACCGAGCTGGGGCGGCCAGCCGAG 344
Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAAACAGCAGCGAGATCTCTGGAGCGAAACCGGGCGGAGGTGGACACGCGCTGC 404
Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db 405 AGACACAACCTACGAGGGCGGAGACCGACCTCTCCCTGCGCGGCTTGAACAGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db 465 GTCCGCATCTCCCTGTCCAGGACGAGCCCTCAACCCACACACACTCTGGTCTGTTCG 524
Qy 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGCCTGCTGGTTCAGGAATGGCCAGGAG 584
Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220

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585 ACAGTGGGGTCTCATCCACACAGCTTTATTAGGATGGGAGCTGGACCTTCCAGGTCCTG 644
221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrCysHisValGluHisPro 240
645 GTCATCTCGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
705 AGCCTGAAGAGCCCAATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
261 GlyGlyGlyGlySer 265
756 GGCGGTGGTGGTTC 770

JLT 13
13963
SEQUENCE 122 from patent US 5869270.
AR033963
AR033963.1 GI:5949568
Unknown.
ORGANISM
Unclassified.
REFERENCE 1 (bases 1 to 1508)
AUTHORS Rhode, P.R., Jiao, J.-A., Burkhardt, M. and Wong, H.C.
TITLE Single chain MHC complexes and uses thereof
JOURNAL Patent: US 5869270-A 122 09-FEB-1999;
FEATURES
Location/Qualifiers
source
1. .1508
/organism="unknown"
/mol_type="unassigned DNA"

Alignment Scores:
i. No.: 2,13e-110 Length: 1508
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 70.7% Indels: 10
Gaps: 3

US-048-116B-6 (1-306) x AR033963 (1-1508)
1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValMetValLeu 20
6 ATGGCTCTGCAGATCCCGAGCCCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTGTGT 65
21 SerSerProGlyThrGluGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCAGCTGAA----- 119
41 IleValValSerLysSerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
120 ATCAACGAAGCTGTCTGTCTAGCGAGGGGGGGAAGC-----GGCGGA 164
61 GlyGlyGlySerGluArgHisPheValValGlnPheGlyGlyGlyCysTyrThrAsn 80
165 GGCGGAACCTCCGAAAGGATTTCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 224
81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
225 GGAGCCAGCGCATACGGCTCGTGTGACCATATCATCAACCGGAGGAGTACGTGGCG 284
101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAlaGlu 120
285 TAGCAGCAGCAGCTGGCGGAGTACCGCGGCTGACCGAGCTGGGGGCGCAGACCGCGAG 344
121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
345 TACTGGAACAGCCGAGCGGAGATCTCTGGAGCGAACCGGGCCGAGGTGGACACGGCGTGC 404
141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160

```

```

405 AGACACACTACGAGGGCGGAGACACCTCCCTCGGCGGCTTCAACAGCCCAAT 464
161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
465 GTGCGCATCTCCCTGTCCAGGAGAGAGGCGCTCAACACCAACACACTCTGGTCTGTCTG 524
181 ValThrAspPheTyrProAlaValIleLysValArgTyrPheArgAsnGlyGlnGluGlu 200
525 GTGACAGATTCTACCCAGCCAGATCAAGTCCGCTGTGTGTGTGTGTGTGTGTGTGTGTGT 584
201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCCAGGTCTCTG 644
221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrCysHisValGluHisPro 240
645 GTCATCTCGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
705 AGCCTGAAGAGCCCAATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
261 GlyGlyGlyGlySer 265
756 GGCGGTGGTGGTTC 770

RESULT 14
AR152030
LOCUS AR152030
DEFINITION Sequence 24 from patent US 6232445.
ACCESSION AR152030
VERSION AR152030.1 GI:15118080
KEYWORDS
SOURCE
ORGANISM
Unclassified.
REFERENCE 1 (bases 1 to 1508)
AUTHORS Rhode, P.R., Acevedo, J., Burkhardt, M., Jiao, J.-A. and Wong, H.C.
TITLE Soluble MHC complexes and methods of use thereof
JOURNAL Patent: US 6232445-A 24 15-MAY-2001;
FEATURES
Location/Qualifiers
source
1. .1508
/organism="unknown"
/mol_type="unassigned DNA"

ORIGIN
Alignment Scores:
Pred. No.: 2,13e-110 Length: 1508
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 70.7% Indels: 10
Gaps: 3

US-10-048-116B-6 (1-306) x AR152030 (1-1508)
1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValMetValLeu 20
6 ATGGCTCTGCAGATCCCGAGCCCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTGTGT 65
21 SerSerProGlyThrGluGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCAGCTGAA----- 119
41 IleValValSerLysSerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
120 ATCAACGAAGCTGTCTGTCTAGCGAGGGGGGGAAGC-----GGCGGA 164
61 GlyGlyGlySerGluArgHisPheValValGlnPheGlyGlyGlyCysTyrThrAsn 80
165 GGCGGAACCTCCGAAAGGATTTCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 224

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GenCore version 5.1.9
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OM protein - nucleic search, using frame_plus_p2n model

Run on: June 30, 2006, 01:23:04 ; Search time 522.277 Seconds
(without alignments)
6007.369 Million cell updates/sec

Title: US-10-048-116B-6_COPY_1_300

Perfect score: 1572

Sequence: 1 MALQIPSLLSAAVVVLVWL.....KKQNAQLKWLKALKKKLAQ 300

Scoring table:

BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 5244920 seqs, 3486124231 residues

Total number of hits satisfying chosen parameters: 10489840

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlp
-Q=/abs/ABSSWEB.spool/US10048116/runat.29062006.093309.10102/app.query.fasta.1
-DB=N Geneseq -QFMT=fastcap -SUFFIX=p2n.rng -MINMATCH=0.1 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blsum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000 -HOST=abs05p
-USER=US10048116 @CGN 1.1423 @runat.29062006.093309.10102 -NCPU=6 -ICPU=3
-NO MMAP -NEG SCORES=0 -WAIT -DSPBLOC=100 -LONGLOG -DEV TIMEOUT=120
-WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N Geneseq 8:*

1: Geneseqn1980s:*
2: Geneseqn1990s:*
3: Geneseqn2000s:*
4: Geneseqn2001as:*
5: Geneseqn2001bs:*
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13: Geneseqn2004bs:*
14: Geneseqn2005s:*
15: Geneseqn2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1572	100.0	921	5 AAF55099	Aaf55099 DNA encod
2	1255.5	79.9	893	2 AAT04262	Aat04262 Hybrid IA
3	1222	77.7	945	12 ADQ31225	Adq31225 I-Ab(beta

	4	1206	76.7	915	12	ADQ31228
5	1161.5	73.9	4724	2	AAV12068	Adq31228 I-Ab(beta
6	1151	73.2	1013	2	AAT04269	Aav12068 Murine IA
7	1145	72.8	1382	2	AAT17588	Aat04269 Hybrid IA
8	1145	72.8	1382	2	AAT86989	Aat17588 Vector SC
9	1145	72.8	1382	8	ACA60744	Aat86989 SCE1 e1ng
10	1145	72.8	1385	2	AAT17586	ACA60744 Mouse MHC
11	1145	72.8	1385	2	AAT86987	Aat17586 Vector SS
12	1145	72.8	1385	8	ACA60742	Aat86987 SSC1 sing
13	1145	72.8	1385	8	AAT17587	ACA60742 Mouse MHC
14	1145	72.8	1508	2	AAT86988	Aat17587 Vector SC
15	1145	72.8	1508	2	AAT89069	Aat86988 SCT1 e1ng
16	1145	72.8	1508	8	ACA60743	Aat89069 Single ch
17	1036.5	69.8	798	12	ADJ75986	ACA60743 Mouse MHC
18	1036.5	69.8	798	14	ADX26090	Adj75986 Marker ge
19	1093	65.5	1085	4	ABI99040	Adx26090 Novel cel
20	1049.5	66.8	1698	4	ABI99038	Abi99040 Murine pC
21	1044.5	66.4	1662	4	ABI99039	Abi99038 Murine pC
22	979.5	62.3	702	2	AAQ03170	Abi99039 Murine pC
23	979.5	62.3	702	2	AAT06286	Aaq03170 Sequence
24	979.5	62.3	702	2	AAQ56920	Aat06286 I-Ab-beta
25	972	61.8	1243	6	ABN84048	Aaq56920 Mouse I-A
26	963.5	61.3	702	2	AAQ35055	Abn84048 Single ch
27	957	60.9	1686	4	ABI99031	Aaq35055 IAB beta
28	957	60.9	1701	4	ABI99028	Abi99031 MBP 1-14
29	957	60.9	2059	4	ABI99032	Abi99028 IAS MBP 1
30	957	60.9	2346	4	ABI99027	Abi99032 MBP 1-14
31	952	60.6	1707	4	ABI99030	Abi99027 IAS MBP 1
32	949	60.4	1680	4	ABI99021	Abi99030 IAS MBP 9
33	949	60.4	2053	4	ABI99029	Abi99021 I-Aa MBP
34	949	60.4	2343	4	ABI99033	Abi99029 IAS MBP 9
35	871	55.4	1323	2	AAT60705	Abi99033 MBP 90-10
36	854.5	54.4	1323	2	AAT60700	Aat60705 CDNA enco
37	844.5	53.7	861	14	AEC64482	Aat60700 CDNA enco
38	839.5	53.4	1192	10	AAD63150	Aec64482 DRB1-biot
39	839.5	53.4	1192	10	AAD62751	Aad63150 Human maj
40	839.5	53.4	1192	11	ADP88246	Aad62751 Human maj
41	839.5	53.4	1192	13	ADR24869	Adp88246 Lung canc
42	833.5	53.0	941	12	ADO40822	Adr24869 Breast ca
43	829	52.7	562	6	ABK63510	Ado40822 DNA encod
44	829	52.7	562	10	ADB57995	Abk63510 Rat seque
45	829	52.7	562	10	ABT41775	Adb57995 Toxicity

ALIGNMENTS

RESULT 1

AAF55099
ID AAF55099 standard; DNA; 921 BP.

AAF55099;

15-MAY-2001 (first entry)

DNA encoding a fusion protein comprising a beta chain of MHC.

Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
major histocompatibility complex; Fc region; antigen; T lymphocyte;
immunostimulant; vaccine; infection; tumour; ss.

Synthetic.

Key	Location/Qualifiers
CDS	1..921
FT	/*tag= a

WO200109194-A1.

08-FEB-2001.

28-JUL-2000; 2000WO-FR002193.

29-JUL-1999; 99FR-00009862.

FT		/product= "IA beta chain beta 2 region"	
FT	primer_bind	521..550	
FT		/*tag= j	
FT		/note= "probable primer binding site (primer #271)"	
FT	primer_bind	532..554	
FT		/*tag= k	
FT		/note= "probable primer binding site (primer #272)"	
FT	primer_bind	808..836	
FT		/*tag= l	
FT		/note= "probable primer binding site (primer #259)"	
FT	primer_bind	877..893	
FT		/*tag= m	
FT		/note= "probable primer binding site (primer #232)"	
XX			
FN	W09523814-A1.		
XX			
XX	08-SEP-1995.		
PD			
XX			
PF	03-MAR-1995;	95WO-US0002689.	
XX			
PR	04-MAR-1994;	94US-00207481.	
XX			
PA	(NAJBE-) NAT JEWISH CENT IMMUNOLOGY & RESPIRATORY.		
XX			
PI	Kappler JW, Marrack P;		
XX			
XX			
DR	WPI; 1995-320543/41.		
DR	P-PSDB; AAR82533.		
XX			
PT	Peptide-MHC complex comprising antigenic peptide, linker and MHC segment		
PT	- useful as reagents for the treatment of diseases including auto-immune		
PT	diseases, immuno-stimulatory diseases or graft-host rejection.		
XX			
PS	Example 1; Page 53; 94pp; English.		
XX			
CC	This sequence represents a hybrid IA beta chain gene, containing the		
CC	chicken ovalbumin peptide (cOVA). This sequence was used in the		
CC	construction of a hybrid IA alpha beta dimer. The encoded protein (prad-		
CC	OVA) was found to be more stable than the IA alpha beta dimer. The		
CC	stability was decreased by the addition of a MHC groove specific binding		
CC	peptide (e.g. see AAR82527, AAR82528 and AAR82531), compared to an		
CC	increase seen on the addition of a MHC binding peptide to IE k/d-MCC.		
CC	These complexes may be used to regulate an immune response. The complexes		
CC	are capable of being recognised by a TCR alone or in combination with		
CC	additional MHC proteins. These complexes are useful for therapeutic		
CC	purposes and experimental purposes. They can also be used as reagents for		
CC	the treatment of diseases including autoimmune diseases, immunodeficiency		
XX	diseases, immunoproliferation diseases, and graft-host rejection		
SQ	Sequence 893 BP; 204 A; 239 C; 275 G; 175 T; 0 U; 0 Other;		
	Alignment Scores:		
	Pred. No.:	9,23e-112	Length: 893
	Score:	1255.50	Matches: 242
	Percent Similarity:	95.4%	Conservative: 6
	Best Local Similarity:	93.1%	Mismatches: 7
	Query Match:	79.9%	Indels: 5
	DB:	2	Gaps: 1
	US-10-048-116B-6_COPY_1_300 (1-300) x AAT04262 (1-893)		
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Db	61	ATGGCTCTGAGATCCCGAGCTCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTG	120
Qy	21	SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro	40
Db	121	AGCAGCCCCGGGACTGAGGCGGAACTCCGTACATGCTGCCCATGCT-----	168
Qy	41	IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly	60
Db	169	--GAGATCAATGAGCTGGCAGAGAGAGTGGGGGCTCCTACTAGTGCCTCGAGGCTCTGGA	225

XX
DR WPI: 2004-546819/53.
DR P-PSDB; ADQ31224.
XX
PT Peptide-Class II major histocompatibility complex (MHC) composite, useful
PT for detecting antigen specific CD4+ T-cell, comprises antigen peptide
PT containing epitope of mucous membrane invasive protein, and extracellular
PT region of MHC.
XX
XX Example 1; SEQ ID NO 10; 30pp; Japanese.

XX The invention relates to a novel class II major histocompatibility
CC complex (MHC) antigenic peptide composite comprising a peptide containing
CC the T-cell antigenic determinant of a mucous membrane invasive protein
CC and the extracellular region of a class II MHC molecule or at least part
CC of the extracellular region of the class II MHC molecule having an amino
CC acid sequence comprising one or more deletions, substitutions or
CC additions. The molecule of the invention may be useful for detecting an
CC antigen-specific CD4+ T-cell by flow cytometry and for presenting a
CC microorganism-derived mucous membrane invasive protein as an antigen. The
CC method of the invention enables efficient detection of antigen-specific
CC activation of CD4+ T-cells in the mucous membrane. The current sequence
CC is that of the class II major histocompatibility complex-related I-
CC Ab(alpha)-Cholera toxin B subunit (CTB)-leucine zipper (LZ)-Bira fusion
CC cDNA of the invention.

XX
SQ Sequence 945 BP; 230 A; 256 C; 294 G; 165 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 1.74e-108 Length: 945
Score: 1222.00 Matches: 242
Percent Similarity: 85.0% Conservative: 14
Best Local Similarity: 80.4% Mismatches: 41
Query Match: 77.7% Indels: 4
DB: 12 Gaps: 3

US-10-048-116B-6_COPY_1_300 (1-300) x ADQ31225 (1-945)

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QY 20 LeuSerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHis 39
DB 64 CTGAGCTCCCACTGGCTTTGGCTGGAGACTCTCGCTGTGGACATAAGACCGCCGAC 123
QY 40 ProlleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySer 59
DB 124 GCGATCGCGCCCATCAGCATGCGCAACGCGAGGTGGTGGTCC---GGTGGAGGGGGAAGT 180
QY 60 GlyGlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrThr 79
DB 181 GGAGGTGGAGGGTCTGAAAGGCAATTCGTGTACCACTTCATGGCGGAGTGCTACTTCACC 240
QY 80 AsnGlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrVal 99
DB 241 AACGGACCGCAGGCATACGATATGTGACAGATACATCTACACCGGGAGGAGTACGTG 300
QY 100 ArgTyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAla 119
DB 301 CGCTACGACAGGACGTGGCGGAGCACCGCGGGTGACCGAGTGGGGCGGCAGACGCC 360
QY 120 GluTyrTrpAsnSerGlnProGluIleLeuGluAaGThrArgAlaGluValAspThrAla 139
DB 361 GAGTACTGGAACAGCCAGCGGAGATCTTGGAGCGAACCGGGCGGAGCTGGACACGGTG 420
QY 140 CysArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnPro 159
DB 421 TGCAGACACAACTACGAGGGCGGAGACCCACACTCTCTCGGGGGCTTGAACAGCCC 480
QY 160 AsnValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHieAsnThrLeuValCys 179
DB 481 AATGTGTCATCTCCCTGTCCAGGACAGAGGGCCCTCAACACAGCACCAACTCTGGTCTGC 540

QY 180 SerValThrAspPheTyrProAlaIleValLeuValArgTrpPheArgAsnGlyGlnGlu 199
DB 541 TCAGTGACAGATTTCTACCCAGCCCAAGATCAAGTCGCTGGTTCGGAAATGCCAGGAG 600
QY 200 GluThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnVal 219
DB 601 GAGCGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTC 660
QY 220 LeuValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHis 239
DB 661 CTGGTCTATCTCGAGATGACCCCTCGCGGGGAGAGGTCTACACCTGTACGTGGAGCAT 720
QY 240 ProSerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSer 259
DB 721 CCCAGCCTGAAGAGCCCCATCTCTGTGGAGTGGAGGCACAGTCGTCAGCA-----GAC 774
QY 260 LysGlyGlyGlySerThrAlaProSerAlaGlnLeuLysLysLysLeuGlnAla 279
DB 775 CTGGTTCGCGCGGATCCACTACAGCTCCATCAGCTCAGTTGAAAAAGAACTGCAGGCA 834
QY 280 LeuLysLysLysAsnAlaGlnLeuLysTrpLysLeuGlnAlaLeuLysLysLysLeuAla 299
DB 835 CTTAAGAAAAAGAACGCTCAGCTGAAGTGGAAACTTCAAGCCCTCAAGAAGAACTCGCC 894
QY 300 Gln 300
DB 895 CAG 897
RESULT 4
ADQ31228
ID ADQ31228 standard; cDNA; 915 BP.
XX
AC ADQ31228;
XX
DT 07-OCT-2004 (first entry)
XX
DE I-Ab(beta)-E. coli heat-labile toxin B subunit-LZ-Bira fusion cDNA.
XX
KW class II major histocompatibility complex; MHC; CD4+ T-cell detection;
KW flow cytometry; mucous membrane invasive antigen;
KW I-Ab(beta)-heat-labile toxin B subunit-leucine zipper-Bira fusion; LTB;
KW ss; gene.
XX
OS Escherichia coli.
OS Unidentified.
XX
FH Key Location/Qualifiers
FT CDS 1..915
FT /tag= a
FT /product= "I-Ab(beta)-Escherichia coli heat-labile toxin
FT B subunit (LTB)-leucine zipper (LZ)-Bira fusion protein"
XX
PN JP2004196789-A.
XX
PD 15-JUL-2004.
XX
PF 03-DEC-2003; 2003JP-00404367.
XX
PR 03-DEC-2002; 2002JP-00351818.
XX
PA (SENT-) SENTAN KAGAKU GIJUTSU INCUBATION CENT KK.
XX
XX WPI: 2004-546819/53.
DR P-PSDB; ADQ31227.
XX
PT Peptide-Class II major histocompatibility complex (MHC) composite, useful
PT for detecting antigen specific CD4+ T-cell, comprises antigen peptide
PT containing epitope of mucous membrane invasive protein, and extracellular
PT region of MHC.
XX
XX Example 3; SEQ ID NO 13; 30pp; Japanese.

CC diabetes, multiple sclerosis, autoimmune thyroiditis, systemic lupus
 CC erythematous, myasthenia gravis, Crohn's disease and inflammatory bowel
 CC disease, or an allergy, e.g. asthma and contact sensitivity
 XX

SQ Sequence 4724 BP; 1196 A; 1194 C; 1200 G; 1134 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 1..11e-101 Length: 4724
 Score: 1161.50 Matches: 228
 Percent Similarity: 85.8% Conservative: 1
 Best Local Similarity: 85.4% Mismatches: 1
 Query Match: 73.9% Indels: 37
 DB: 2 Gaps: 2

US-10-048-116B-6_COPY_1_300 (1-300) x RAV12068 (1-4724)

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 |||||
 QY 21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
 |||||
 Db 511 AGAGCCCGAGGGACTGAGGCGGAAC----- 537
 |||||
 QY 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 |||||
 Db 537 ----- 537
 |||||
 QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTrpThrAsn 80
 |||||
 Db 538 -----TCCGAAGGATTCGTGGTCCAGTTCAGGGCGAGTCTACTACCAAC 588
 |||||
 QY 81 GlyThrGlnArgIleArgLeuValThrArgTrpIleTyrAsnArgGluGluTrpValArg 100
 |||||
 Db 589 GGGACGACGGCATACGGCTCGTGACCATATACACCGGGAGGAGTACGGTGC 648
 |||||
 QY 101 TyrAspSerAspValGlyGluTrpArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 |||||
 Db 649 TACGACAGCGAGCTGGCGGAGTACCGCGGTGACCGAGCTGGGGGGCGCAGCGCGAG 708
 |||||
 QY 121 TyrTrpAsnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 |||||
 Db 709 TACTGGAACAGCGAGCGGAGATCTCTGGAGCGAACCGGGCGGAGTGGACACGGCGTCC 768
 |||||
 QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGlnProAsn 160
 |||||
 Db 769 AGACACAACTACAGAGGGCGGGAGACAGACCTCTCTGGCGGCTTGAACAGGCCAAT 828
 |||||
 QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
 |||||
 Db 829 ATCGCCATCTCCCTGTCCAGGACAGAGGCCCTCAACCAACCAACACTCTGGTCTGTTCG 888
 |||||
 QY 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
 |||||
 Db 889 GTGACAGATTTCTACCCAGCAGAGATCAAAAGTGCCTGTTCAGGAATGCCAGGAGGAG 948
 |||||
 QY 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220
 |||||
 Db 949 ACAGTGGGGGTCTCATCCACACAGCTATTAGGAATGGGACTGGACCTTCCAGGTCCTG 1008
 |||||
 QY 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 |||||
 Db 1009 GTCATGCTGGAGATGACCCCTCATCAGGGAGAGGTTCTACCTGCCATGTGGAGCATCCC 1068
 |||||
 QY 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGlnSerAlaArgSerLys 260
 |||||
 Db 1069 AGCCTGAAGAGCCCCATCACTGTGGAGTGGAGGGCACAGTCCGAGTCTCCCGGAGCAAG 1128
 |||||
 QY 261 -----GlyGlyGlyGly 264
 |||||
 Db 1129 ATGTTGAGCGGCATCGGGGCG 1149

RESULT 6

AAT04269
 ID AAT04269 standard; DNA; 1013 BP.
 XX
 AC AAT04269;
 XX
 DT 16-APR-1996 (first entry)
 XX
 DE Hybrid IA beta chain gene.
 XX
 KW Major histocompatibility complex; MHC; T-cell receptor; TCR;
 KW autoimmune disease; immunodeficiency disease; immune response;
 KW immunoproliferation disease; graft-host rejection; therapy; B cell;
 KW M12.C3; pM12-1Ab-Ea; ss.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT primer_bind 1..18
 FT /tag= a
 FT /note= "probable primer binding site (primer #76)"
 FT complement(40..74)
 FT primer_bind
 FT /tag= b
 FT /note= "binding site for primer #362 (see AAT04270)"
 FT CDS
 FT 63..959
 FT /tag= c
 FT /product= "hybrid IA beta chain"
 FT 63..143
 FT /tag= d
 FT /note= "leader region"
 FT complement(140..191)
 FT primer_bind
 FT /tag= e
 FT /note= "binding site for primer #363 (see AAT04271)"
 FT complement(177..226)
 FT primer_bind
 FT /tag= f
 FT /note= "primer #364 binding site"
 FT complement(212..266)
 FT primer_bind
 FT /tag= g
 FT /note= "primer #365 (see AAT04272) binding site"
 FT 385..403
 FT /tag= h
 FT /note= "probable primer binding site (primer #270)"
 FT 531..959
 FT /tag= i
 FT /product= "IA beta chain beta 2 region"
 FT 535..564
 FT /tag= j
 FT /note= "probable primer binding site (primer #271)"
 FT 544..568
 FT /tag= k
 FT /note= "probable primer binding site (primer #272)"
 FT 823..850
 FT /tag= l
 FT /note= "probable primer binding site (primer #259)"
 FT 942..976
 FT /tag= m
 FT /note= "probable primer binding site (primer #366)"
 FT 1000..1013
 FT /tag= n
 FT /note= "probable primer binding site (primer #59)"
 FT
 XX WO9523814-A1.
 PN
 XX
 PD 08-SEP-1995.
 XX
 PF 03-MAR-1995; 95WO-US002689.
 XX
 PR 04-MAR-1994; 94US-00207481.
 XX
 PA (NAJE-) NAT JEWISH CENT IMMUNOLOGY & RESPIRATORY.
 XX
 PI Kappler JW, Marrack P;
 XX
 DR WPI; 1995-320543/41.

DR P-PSDB; AAR82538.

XX Peptide-MHC complex comprising antigenic peptide, linker and MHC segment
 PT - useful as reagents for the treatment of diseases including auto-immune
 PT diseases, immuno-stimulatory diseases or graft-host rejection.

XX Example 2; Page 65; 94pp; English.

XX This sequence represents a hybrid IA beta chain gene. This sequence
 CC contains a fragment of the IE alpha chain (residues 56-73), as well as a
 CC linker and cleavage site. This sequence was transfected into a B cell
 CC line (M12.C3) using plasmid pM12-iAb-Ea. It was found that the encoded
 CC sequence was expressed in these cells. Complexes such as this may be used
 CC to regulate an immune response. The complexes are capable of being
 CC recognised by a TCR alone or in combination with additional MHC proteins.
 CC These complexes are useful for therapeutic purposes and experimental
 CC purposes. They can also be used as reagents for the treatment of diseases
 CC including autoimmune diseases, immunodeficiency diseases,
 CC immunoproliferation diseases, and graft-host rejection

SQ Sequence 1013 BP; 220 A; 272 C; 327 G; 192 T; 0 U; 2 Other;

Alignment Scores:

Pred. No.: 1.43e-101 Length: 1013
 Score: 1151.00 Matches: 230
 Percent Similarity: 86.8% Conservative: 6
 Best Local Similarity: 84.6% Mismatches: 22
 Query Match: 73.2% Indels: 14
 DB: 2 Gaps: 4

US-10-048-116B-6_COPY_1_300 (1-300) x AAT04269 (1-1013)

QY 1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db 63 ATGGCTCTGCAGATCCACGCTCTCTCTCGCTGCTGCTGCTGCTGCTGCTGCTGCTG 122
 QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 123 AGCAGCCAGGGAGTGGAGGCTCTGAAAGCATTTCTGTACCATCTTCATCGGCGAG 173
 QY 41 -----11eValValSerGlySerTrpAspGlyGlyGlyGlySerLeuVal 55
 Db 174 GTGCTACTGCCAACATGCTGTGCAACAGCTGGAGGTGGTGTGATCCGGTGA----- 227
 QY 56 ProArgGlySerGlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlu 75
 Db 228 ---GGGGAGAGTGGAGGTGGAGGCTCTGAAAGCATTTCTGTACCATCTTCATCGGCGAG 284
 QY 76 CysTyrTyrThrAsnGlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArg 95
 Db 285 TGCTACTTCCACCAACCGGACGCGCATACGATATGTGACCATACATCTACACCGG 344
 QY 96 GluGluTyrValArgTyrAspSerAspValGlyGluThrArgAlaValThrGluLeuGly 115
 Db 345 GAGAGTACTCGCTACGACAGCGAGCTGGGCGAGCCCGCGGTGACCGAGCTGGGG 404
 QY 116 ArgProAspAlaGluTyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGlu 135
 Db 405 CGGCCACAGCCCGAGTACTGGAACAGCCAGCCGAGATCTCTGGAGCGAAGCGCGCGAG 464
 QY 136 ValAspThrAlaCysArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArg 155
 Db 465 GTGGACACGGGTGTCAGACACAACACTACGAGGGGCCGAGAGCCACACCTCCCTGCGCGG 524
 QY 156 LeuGluGlnProAsnValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsn 175
 Db 525 CTTGAACAGCCCAATGTGTCATCTCTCTGTCAGGACAGAGCCCTCAACACCAAC 584
 QY 176 ThrLeuValCysSerValThrAspPheTyrProAlaLysIleLysValArgTrpPheArg 195
 Db 585 ACTCTGTCTGCTCAGTGACAGATTTCTACCCAGCCAGATCAAGTGGCTGCTGCTCCGG 644
 QY 196 AsnGlyGlnGluThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrp 215

Db 645 AATGCCAGGAGGAGCGGTGGGCTCTCATCCACACAGCTTATTAGGAATGGGACTGG 704
 QY 216 ThrPheGlnValLeuValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCys 235
 Db 705 ACCTTCAGAGTCTGCTGCTCATGCTGGAGATGACCCCTCGGCGGGGAGAGGTTTAVACCTGT 764
 QY 236 HisValGluHisProSerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGlu 255
 Db 765 CACGTGAGCATCCAGCCTGAGAGGCCCATCATCTGTGGAGTGGAGGCGACAGTCTCGAG 824
 QY 256 SerAlaArgSerLys-----GlyGlyGlyGly 264
 Db 825 TCTGCTGGAGCAAGATGTTGAGCGGCATCGGGGCG 860
 RESULT 7
 AAT17588
 ID AAT17588 standard; DNA; 1382 BP.
 XX
 AC AAT17588;
 XX
 DT 26-SEP-1996 (first entry)
 XX
 DE Vector SCE1-derived single chain gene encoding MHC fusion complex.
 XX
 KW MHC; major histocompatibility complex; PCR; polymerase chain reaction;
 KW T cell activity modulator; antagonist; immune disorder; allergy;
 KW multiple sclerosis; insulin-dependent diabetes mellitus;
 KW rheumatoid arthritis; myasthenia gravis; ds.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT CDS 6..1382
 FT /*tag= a
 FT sig_peptide 6..86
 FT /*tag= b
 FT /label= I-Ad beta_chain_leader
 FT /note= "murine MHC class II I-Ad gene beta chain leader
 FT sequence"
 FT misc_feature 87..137
 FT /*tag= c
 FT /label= OVA_323-339
 FT /note= "chicken ovalbumin residues 323-339"
 FT misc_feature 138..167
 FT /*tag= d
 FT /note= "10 residue linker peptide"
 FT misc_feature 168..452
 FT /*tag= e
 FT /label= I-Ad beta1
 FT /note= "murine MHC class II I-Ad gene beta-1 domain"
 FT misc_feature 453..734
 FT /*tag= f
 FT /label= I-Ad beta2
 FT /note= "murine MHC class II I-Ad gene beta-2 domain"
 FT misc_feature 735..806
 FT /*tag= g
 FT /note= "24 residue peptide linker"
 FT misc_feature 807..1067
 FT /*tag= h
 FT /label= I-Ad alpha1
 FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
 FT misc_feature 1068..1352
 FT /*tag= i
 FT /label= I-Ad alpha2
 FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
 FT misc_feature 1353..1379
 FT /*tag= j
 FT /note= "EE tag"
 XX
 PN WO9604314-A1.
 XX
 XX 15-FEB-1996.

US-10-048-116B-6_COPY_1_300 (1-300) x AAT86987 (1-1385)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 6 ATGGCTTCGACAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTG 65
 Qy 21 SerSerProGlyThrGluGlyGlyValSerIleCysPheSerProSerLeuGluHisPro 40
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTCAAGGCGAGTGTCTACGCTGAA----- 119
 Qy 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 120 ATCAACGAGCTGTGTCTAGCGGAGGGGGCGGAGC-----GGCGGA 164
 Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTrpTrpThrAsn 80
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 165 GGGGGAACCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTGTCTACTACACCAAC 224
 Qy 81 GlyThrGlnArgIleArgLeuValThrArgTrpIleTyrAsnArgGluGluTyrValArg 100
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 225 GGGACGACGACGATACCGCTCGTACCGAGATACATCTACCAACCGGGAGGAGTACGTGCGC 284
 Qy 101 TyrAspSerAspValGlyGlyValThrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 285 TACGACGACGACGTGGGGAGTACCGCGGTGACCGAGTGGGGCGGCCAGACGCCGAG 344
 Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 345 TACTGGAACACCGACCGGAGATCTTGAGCGAAGCGGGCGGAGTGGACACGGCGTGC 404
 Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 405 AGACACAACTACGAGGGGCGGAGACCGACCTCTCTGCGGCGGTGTGAACACGCCCAAT 464
 Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer 180
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 465 GTCCGCATCTCCCTGTCCAGCAGAGAGCCCTCAACCAACACACATCTGTGTGTGTG 524
 Qy 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 525 GTGACAGATTCTACCCAGCCAGATCAAGTGGCGCTGTTCAGGAATGGCCAGGAGGAG 584
 Qy 201 ThrValGlyValSerSerThrGlnIleArgAsnGlyAspTrpPheGlnValLeu 220
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 585 ACAGTGGGGGTCTCATCCACACAGCTATTAGGAATGGGGAGTGGACCTTCCAGGTCTGT 644
 Qy 221 ValMetLeuGluMetThrProHisGlnGlyGlyValTyrThrCysHisValGluHisPro 240
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 645 GTCATGTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
 Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
 Qy 261 GlyGlyGlyGlySer 265
 Db | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
 756 GCGCGTGTGTGTCTCC 770

RESULT 12

ACA60742

ID ACA60742 standard; DNA; 1385 BP.

AC ACA60742;

XX ACA60742;

DT 16-JUN-2003 (first entry)

DE Mouse MHC I-Ad/Ova 323-339 synthetic gene SSC1.

XX MHC; major histocompatibility complex; gene therapy; fusion complex;
 KW peptide-binding groove; T cell modulation; class II MHC; vaccine;
 KW autoimmune disorder; multiple sclerosis; rheumatoid arthritis;
 KW insulin-dependent diabetes mellitus; myasthenia gravis; immunogen;

Chronic allergy; mouse; ds; I-Ad; gene.

Mus sp.

Synthetic.

US2002198144-A1.

26-DEC-2002.

06-JUL-2001; 2001US-00900379.

29-JUL-1994; 94US-00283302.

01-FEB-1995; 95US-00382454.

17-JAN-1997; 97US-00776084.

(DADE-) DADE INT INC.

Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;

Chavallaz P, Jiao JJJ;

WPI; 2003-341126/32.

P-PSDB; ABU72106.

Novel major histocompatibility complex fusion complex having presenting peptide covalently linked to MHC molecule containing peptide-binding groove, used for suppressing immune response in multiple sclerosis, allergies.

Example 17; Fig 27; 126pp; English.

The invention relates to a major histocompatibility complex (MHC) fusion complex (I) comprising an MHC molecule that contains a peptide-binding groove, and a presenting peptide covalently (e.g. an antigenic peptide) linked to the MHC molecule, where (I) is capable of modulating the activity of a T cell. Also included are a DNA construct coding for the complex, where the MHC molecule is a class II MHC (e.g. mouse I-Ad or I-As, or human HLA-DR1 (human leukocyte antigen-DR1)), a multivalent MHC fusion complex comprising two or more linked complexes, identifying a peptide that can modulate the activity of T cells (involving introducing into host cells cloning vectors that each contain the fusion complex DNA, culturing the host cells under conditions suitable for expression of the MHC fusion complex, and selecting host cells that express MHC fusion complex that modulate the activity of T cells), a single recombinant expression vector comprising DNA that codes for the alpha and beta chains of the fusion complex MHC protein, a single recombinant expression vector comprising DNA that codes for a T cell costimulatory factor and the alpha and beta chains of the MHC fusion complex. The DNA constructs can contain heterologous leader peptide sequences and Kozak sequence for efficient expression of the fusion complex. Also included are inducing an immune response in a mammal (including vaccinating a mammal against a targeted disorder, by administering DNA sequence comprising a fusion complex, or DNA sequence coding for a fusion complex which is a single chain fusion molecule) and suppressing an immune response in a mammal by administering to the mammal a DNA sequence comprising an expression vector, encoding a full length MHC molecule that contains a transmembrane domain, and a presenting peptide that is a T cell receptor (TCR) antagonist or partial agonist and is covalently linked to the MHC protein, or DNA sequence coding for the fusion complex which is a single chain fusion molecule. The methods are useful for identifying a peptide that can modulate the activity of T cells, inducing an immune response in a mammal (including vaccinating a mammal against a targeted disorder) and for suppressing an immune response in a mammal. The disorders include an autoimmune disorder such as multiple sclerosis, insulin-dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or chronic allergies. The present sequence encodes a mouse MHC class II I-Ad fusion complex of the invention

SQ Sequence 1385 BP; 316 A; 383 C; 399 G; 287 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 8.33e-101 Length: 1385

Score: 1145.00 Matches: 227

Percent Similarity: 87.2% Conservative: 4

Best Local Similarity: 85.7%
 Query Match: 72.8%
 DB: 8 Mismatches: 24
 10
 3 Gaps:

US-10-048-116B-6_COPY_1_300 (1-300) x ACA60742 (1-1385)

QY 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db 6 ATGGCTCTGAGATCCCGACCTCTCTCAGCTGTGTGGTGGTGTGGTGTGGTGTG 65
 QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGCAGCCAGGACCTTAAGTATCTCTCAGGCTGTTCAGCTGTTCAGCTGTTCAG 119
 QY 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db 120 ATCAACGAAGCTGT 164
 QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheGlyGlyGlyCysTyrThrAsn 80
 Db 165 GGGGGAACCTCCGAAGGCAATTCGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 224
 QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 Db 225 GGGACCCAGCGCATACGGCTCTGACAGATACATCTACACCGGAGGAGTACGTGGC 284
 QY 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db 285 TACGACAGCGAGCTGGCGGAGTACCGCGGTGTGTGTGTGTGTGTGTGTGTGTGTGT 344
 QY 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 Db 345 TACTGGAACAGCGAGCGCGAGATCTGTGAGCGAAGCGCGCGGTGTGTGTGTGTGTGT 404
 QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
 Db 405 AGACACAACTACGAGGGCGGAGCAGCAGCCTCTCTGGCGGCTTGAACAGCCCAAT 464
 QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
 Db 465 GTCCCATCTCCCTGTCCAGGACAGAGGCGCTCAACCAACCAACCACTCTGTGTGTGT 524
 QY 181 ValThrAspPheTyrProAlaValIleValValArgTyrPheArgAsnGlyGlnGluGlu 200
 Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAAGTGTGTGTGTGTGTGTGTGTGTGTGT 584
 QY 201 ThrValGlyValSerThrGlnLeuIleArgAsnGlyAspTyrPheGlnValLeu 220
 Db 585 ACAGTGGGGTCTCATCCACAGCTTATAGGATGGGACTGGACCTTCCAGGTCTCTG 644
 QY 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 Db 645 GTCATGCTGGAGATGACCTCTCATCAGGAGAGGTCTACACCTGCCATGTGGGAGCATCC 704
 QY 241 SerLeuLysSerProIleThrValGluThrArgAlaGlnSerGluSerAlaArgSerLys 260
 Db 705 AGCCTGAAGAGGCCCATCACTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 755
 QY 261 GlyGlyGlyGlySer 265
 Db 756 GCGCGTGTGTGTTC 770

RESULT 13

AAT17587
 ID AAT17587 standard; DNA; 1508 BP.
 XX
 AC AAT17587;
 XX
 DT 26-SEP-1996 (first entry)
 XX
 DE Vector SCTL-derived single chain gene encoding MHC fusion complex.
 XX
 KW MHC; major histocompatibility complex; PCR; polymerase chain reaction;

KW T cell activity modulator; antagonist; immune disorder; allergy;
 KW multiple sclerosis; insulin-dependent diabetes mellitus;
 KW rheumatoid arthritis; myasthenia gravis; da.
 XX
 XX Synthetic.

Key Location/Qualifiers
 CDS 6..1508
 /tag= a
 sig_peptide 6..86
 /tag= b
 /label= I-Ad beta chain_leader
 /note= "murine MHC class II I-Ad gene beta chain leader sequence"
 misc_feature 87..137
 /tag= c
 /label= OVA 323-339
 /note= "chicken ovalbumin residues 323-339"
 misc_feature 138..167
 /tag= d
 /note= "10 residue linker peptide"
 misc_feature 168..452
 /tag= e
 /label= I-Ad beta1
 /note= "murine MHC class II I-Ad gene beta-1 domain"
 misc_feature 453..734
 /tag= f
 /label= I-Ad beta2
 /note= "murine MHC class II I-Ad gene beta-2 domain"
 misc_feature 735..806
 /tag= g
 /note= "24 residue peptide linker"
 misc_feature 807..1067
 /tag= h
 /label= I-Ad alpha1
 /note= "murine MHC class II I-Ad gene alpha-1 domain"
 misc_feature 1068..1352
 /tag= i
 /label= I-Ad alpha2
 /note= "murine MHC class II I-Ad gene alpha-2 domain"
 misc_feature 1353..1505
 /tag= j
 /label= I-Ad alpha-TM
 /note= "murine MHC class II I-Ad gene alpha-transmembrane domain"

W09604314-A1.

15-FEB-1996.

31-JUL-1995; 95WO-US009816.

29-JUL-1994; 94US-00283302.

01-FEB-1995; 95US-00382454.

(DADE-) DADE INT INC.

Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
 Chavallaz P, Jiao J;

WPI; 1996-129343/13.

P-PSDB; AAR98906.

Major histocompatibility complex fusion complex for modulating T cell activity - used in the treatment of immune disorders, e.g. multiple sclerosis, IDDM and rheumatoid arthritis.

Example 17; Fig 28; 210pp; English.

AAT17587 encodes a murine MHC fusion complex capable of modulating T cell activity encoded by the vector SCTL. The MHC fusion complex comprises at least one MHC molecule containing a peptide-binding groove and a presenting peptide covalently linked to the MHC molecule and opt. a

CC transmembrane domain. DNA encoding a MHC fusion complex may be cloned
 CC into a host cell to express the complex. The transformed cells may then
 CC be used to identify peptides that modulate, pref. antagonise, T cell
 CC activity. DNA encoding a MHC fusion complex or a single chain fusion
 CC molecule may be used to vaccinate a mammal against a targeted disorder.
 CC The fusion complexes may be used to suppress an immune response in an
 CC animal suffering from an immune disorder e.g. multiple sclerosis, insulin
 CC -dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or
 CC chronic allergies. The complexes may also be used in the treatment of
 CC livestock and pets such as cats and dogs. The MHC fusion complexes can be
 CC produced such that they contain a single antigenic peptide including one
 CC of known structure, additionally a wide range of peptides can be
 CC presented for T cell interaction

XX
 SQ Sequence 1508 BP; 337 A; 414 C; 440 G; 317 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 9.35e-101 Length: 1508
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 72.8% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116B-6_COPY_1_300 (1-300) x AAT17587 (1-1508)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGCTGATGCTGCTG 65
 Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGCTGTTCAGCTGTTCACGCTGAA----- 119
 Qy 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db 120 ATCAACGAGCTGCTGCTAGCGAGGGGGGGGAGC-----GGCGGA 164
 Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheIleGlyGlyGlyGlyGlyGly 80
 Db 165 GGGGGAACCTCCGAAGGCAATTCGTGTGCTAGCGAGGGGGGGGAGC----- 224
 Qy 81 GlyThrGlnArgIleArgLeuValThrArgTrpIleTyrAsnArgGluGluTyrValArg 100
 Db 225 GGGAGCGAGCGATACGGCTCGTGACCAAGATACATCAACCGGGAGGAGTACGTGCGC 284
 Qy 101 TyrAspSerAspValGlyGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db 285 TACGACGCGAGCTGGCGAGTACCGCGGTGACCGAGCTCGGGCGCCAGACGCCGAG 344
 Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 Db 345 TACTGGAACAGCAGCCGAGATCTCTGGAGCGAACCGGGCGGAGGTGGACACGCGCTGC 404
 Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgGluGluGlnProAsn 160
 Db 405 AGACAACTACGAGGGCGGAGACCAACACCTCCCTCGCGCGCTTGAACAGCCCAAT 464
 Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
 Db 465 GTCGCCATCTCCCTCTCCAGGACAGAGCCCTCAACACCAACACACACTCTGCTGTTCG 524
 Qy 181 ValThrAspPheTyrProAlaIleIleIleValArgTrpPheArgAsnGlyGlnGluGlu 200
 Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGCCTGTTTTCAGGAATGGCCAGGAGAG 584
 Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220
 Db 585 ACATGGGGGCTCATCCACACAGCTATTATAGGAATGGGAGCTGGACCTTCCAGTCTCTG 644
 Qy 221 ValMetLeuGluMetThrProHisGlnGlyValTyrThrCysHisValGluHisPro 240
 Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704

Qy 241 SerLeuIysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerIys 260
 Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
 Qy 261 GlyGlyGlyGlySer 265
 Db 756 GGCGGTGGTGGTTCC 770

RESULT 14

AAT86988
 ID AAT86988 standard; DNA; 1508 BP.

AC AAT86988;

XX 27-MAR-1998 (first entry)

XX SCT1 single chain gene.

XX Construction; major histocompatibility complex; MHC; fusion complex;

KW SCT1 single chain gene; sb.

XX Synthetic.

XX Key Location/Qualifiers

FT CDS 6..1508

FT /*tag= a

XX WO9728191-A1.

XX 07-AUG-1997.

XX 30-JAN-1997; 97WO-US001617.

XX 31-JAN-1996; 96US-00596387.

XX (DADE-) DADE INT INC.

XX Rhode PR, Jiao J, Burkhardt M, Wong HC;

XX WPI; 1997-402555/37.

XX P-PSDB; AAW29213.

XX Single chain major histocompatibility complex comprising linked alpha and
 PT beta chains - useful for suppressing an immune response to an auto-immune
 PT disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes
 PT mellitus, etc.

XX Example 17; Page 137-139; 217pp; English.

XX The present sequence was used in the construction of major

CC histocompatibility complex (MHC) fusion complexes

XX Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 9.35e-101 Length: 1508
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 72.8% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116B-6_COPY_1_300 (1-300) x AAT86988 (1-1508)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20

Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGTGTGCTGCTGATGCTGCTG 65

Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40

Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGCTGTTCAGCTGTTCACGCTGAA----- 119

QY 41 IleValValSerClySerTrpaspGlyGlyGlySerLeuValProargGlySerGly 60
Db 120 ATCAACGAAGCTGCTCGTACGAGGGGGCGGAAGC-----GGCGGA 164
QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrThrAsn 80
Db 165 GGGGAAACTCCGAAAGGCATTCGTGGTCCAGTTCAGGGCGAGTGCTACTACACCAAC 224
QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrTrpIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGACGACGCGCATACGGCTCGTGACAGATACATCTACACCGGAGGAGTACGTGCGC 284
QY 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProaspAlaGlu 120
Db 285 TAGCAGCAGCGAGTGGCGGAGTACCGCGGGTGACCGAGCTGGGGCGGCAGACGCCGAG 344
QY 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAAACGACGCGGAGATCTTGAGCGNACCGGGCGGAGGTGGACACGGCGTGC 404
QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgLeuGluGlnProAsn 160
Db 405 AGACACAACCTACGAGGGCGGAGACACGACCTCCCTCGCGCGGCTTGAACAGCCCAAT 464
QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer 180
Db 465 GTGCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACCCACCAACACTCTGGTCTGTTCG 524
QY 181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGluGlu 200
Db 525 GTGACAGATTTTACCACCCAGCATCAAGTGGCTGTTTCCAGGATGGCCAGGAGGAG 584
QY 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
Db 585 ACAGTGGGGGTCTCATCCACACAGACTTATTAGGAATGGGAGCTGGACCTTCCAGGTCCG 644
QY 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATGCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCC 704
QY 241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTG-----ACTAGTGGTGGCGGTGGCAGC 755
QY 261 GlyGlyGlyGlySer 265
Db 756 GCGGTGGTGGTTC 770
RESULT 15
AAx89069
ID AAX89069 standard; DNA; 1508 BP.
XX AC AAX89069;
XX AC AAX89069;
DT 14-SEP-1999 (first entry)
DE Single chain IAD/OVA 323-229 MHC fusion protein encoding DNA.
XX Major histocompatibility complex; MHC; single chain MHC; sc-MHC; Ig;
KW peptide binding groove; immunoglobulin; T cell receptor; immune response;
KW immune-related disorder; antigenic peptide; fusion protein; ss.
XX Synthetic.
XX OS
XX PN W09921572-A1.
XX PD 06-MAY-1999.
XX PF 13-OCT-1998; 98WO-US021520.
XX PR 29-OCT-1997; 97US-00960190.
XX PA (SUNO-) SUNOL MOLECULAR CORP.

XX Rhode PR, Acevedo J, Burkhardt M, Jiao J, Wong HC;
XX WPI; 1999-418411/35.
DR P-PSDB; AAV27111.
XX Single chain major histocompatibility complex class I complexes.
PT
XX
XX Example 1; Fig 1; 148pp; English.
XX The invention relates to new single chain major histocompatibility
CC complex (sc-MHC) class II complexes that comprise a peptide binding
CC groove, and a modified class II beta 2 chain or covalently linked
CC immunoglobulin (Ig) light chain constant (CI) region. The MHC complexes
CC are useful for detection and analysis of peptide ligands, pathogenic T-
CC cells, for functional, cellular and molecular assays. They can be used to
CC identify and isolate T cell receptor and/or MHC agonists and antagonists.
CC They can be used in vivo to compete with pathogenic antigen presenting
CC cells involved in immune-related disorders. They can also be used to
CC raise antibodies and to screen immune cells. It is also use in a method
CC of suppressing an immune response in mammals. The sc-MHC complexes
CC comprising modified class II beta 2 chains and/or Ig-CI regions are
CC soluble and provide enhanced yield. These MHC complexes also can contain
CC single antigenic peptides readily isolated from expressing cells in
CC significant quantities. The polyspecific MHC complexes also provide a
CC means to detect cells expressing multiple target structures with a single
CC complex. The present sequence represents a DNA encoding a single chain
CC IAD/OVA 323-229 MHC fusion protein
XX
SQ Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;
Alignment Scores:
Pred. No.: 9,35e-101 Length: 1508
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservatve: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 72.8% Indels: 10
DB: 3 Gaps: 3
US-10-048-116B-6_COPY_1_300 (1-300) x AAX89069 (1-1508)
QY 1 MetalAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTGT 65
QY 21 SerSerProGlyThrGluGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAAGCTGCTCAGCTGAA----- 119
QY 41 IleValValSerGlySerTrpaspGlyGlyGlySerLeuValProargGlySerGly 60
Db 120 ATCAACGAAGCTGCTCGTGTAGCGAGGGGGCGGAAGC-----GGCGGA 164
QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrThrAsn 80
Db 165 GGGGAAACTCCGAAAGGCATTCGTGGTCCAGTTCAGGGCGAGTGCTACTACACCAAC 224
QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrTrpIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGACGACGCGCATACGGCTCGTGACAGATACATCTACACCGGAGGAGTACGTGCGC 284
QY 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProaspAlaGlu 120
Db 285 TAGCAGCAGCGAGTGGCGGAGTACCGCGGGTGACCGAGCTGGGGCGGCAGACGCCGAG 344
QY 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAAACGACGCGGAGATCTTGAGCGNACCGGGCGGAGGTGGACACGGCGTGC 404
QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgLeuGluGlnProAsn 160
Db 405 AGACACAACCTACGAGGGCGGAGACACGACCTCCCTCGCGCGGCTTGAACAGCCCAAT 464


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Qy 161 ValAlaLeSerLeuSerArgThrGluAlaLeuAenHisHisAenThrLeuValCysSer 180
Db 465 GTCCGCATCTCCCTGTCCAGGACAGAGCCCTCAACCAACCAACACTCTGGTCTGTTCG 524
Qy 181 ValThrAspPheTyrProAlaValIleValArgTrpPheArgAenGlyGlnGluGlu 200
Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAAAGTGCCTGGTTCAGGAATGGCCAGAGGAG 584
Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAenGlyAspTrpThrPheGlnValLeu 220
Db 585 ACAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTCTG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy 241 SerLeuIysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerIys 260
Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGGGTGGTGGTTCC 770
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Job time : 528.277 secs

GenCore version 5.1.9
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OM nucleic - nucleic search, using sw model

Run on: June 29, 2006, 23:14:09 ; Search time 5748 Seconds
(without alignments)
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Title: US-10-048-116B-7
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Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 6366136 seqs, 31973710525 residues

Total number of hits satisfying chosen parameters: 12732272

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 4: gb_pl.*
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- 7: gb_sts.*
- 8: gb_sy.*
- 9: gb_un.*
- 10: gb_vi.*
- 11: gb_ov.*
- 12: gb_htg.*
- 13: gb_ini.*
- 14: gb_om.*
- 15: gb_ba.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	918	100.0	921	2	AX081281 Sequence
2	696.2	75.8	893	2	AR047947 Sequence
3	637.6	69.5	1013	2	AR047957 Sequence
4	607.8	66.2	1382	2	AR033964 Sequence
5	607.8	66.2	1382	2	AR175097 Sequence
6	607.8	66.2	1382	2	CS079301 Sequence
7	607.8	66.2	1382	2	AX032545 Sequence
8	607.8	66.2	1385	2	AR033962 Sequence
9	607.8	66.2	1385	2	AR175095 Sequence
10	607.8	66.2	1385	2	CS079299 Sequence
11	607.8	66.2	1385	2	AX032543 Sequence
12	607.8	66.2	1508	2	AR033963 Sequence
13	607.8	66.2	1508	2	AR152030 Sequence
14	607.8	66.2	1508	2	AR175096 Sequence
15	607.8	66.2	1508	2	CS079300 Sequence
16	607.8	66.2	1508	2	AX032544 Sequence
17	604.6	65.9	1508	2	BD138632 Soluble M
18	598.6	65.2	4724	2	AR199666 Sequence

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27	541.4	59.0	1204	6	BC008168	BC008168 Mus muscu
28	539.6	58.8	578	6	AY303784	AY303784 Mus muscu
29	538.2	58.6	575	6	AY303785	AY303785 Mus muscu
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33	524	57.1	729	6	AF293059	AF293059 Mus muscu
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35	522.2	56.9	777	6	AF065913	AF065913 Mus muscu
36	522.2	56.9	792	6	AF293060	AF293060 Mus muscu
37	522.2	56.9	792	6	MUSMHIABS	M13400 Mouse MHC c
38	522.2	56.9	792	6	MUSMHIH2	M66213 Mouse MHC c
39	522.2	56.9	1162	6	BC057998	BC057998 Mus muscu
40	520	56.6	728	6	AF015282	AF015282 Mus muscu
41	519	56.5	760	6	MUSMHIABF	M13541 Mouse MHC c
42	519	56.5	1078	6	AF293061	AF293061 Mus muscu
43	517.4	56.4	792	6	MUSMHIABK	M13538 Mouse MHC c
44	517.4	56.3	725	6	AF293058	AF293058 Mus muscu
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ALIGNMENTS

RESULT 1	AX081281	AX081281	921 bp	DNA	linear	PAT 27-FEB-2001
LOCUS	Sequence 2 from Patent WO0109194.					
DEFINITION	AX081281					
ACCESSION	AX081281					
VERSION	AX081281.1	GI:13170131				
KEYWORDS	synthetic construct					
SOURCE	other sequences; artificial sequences.					
ORGANISM	1					
REFERENCE	Glaichenhaus, N. and Malherbe, L.					
AUTHORS	Recombinant proteins and molecular complexes derived therefrom,					
TITLE	analogous to molecules involved in immune responses					
JOURNAL	Patent: WO 0109194-A 2 08-FEB-2001;					
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	VAISLRYEALNHNHTLVCSVTDFYPAKIKVWRNQGQETVGSSTQILRNDMTFQ					
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	VLVLMETPHGEVYVTCHEHPSLKSPITVEMRAQSESARSKGGGSGTAPSAQLKK					
	LQALKKNAQLKWLQALKLQAQHSHHHH"					
ORIGIN						
	Query Match	100.0%	Score 918;	DB 2;	Length 921;	
	Best Local Similarity	100.0%	Pred. No. 4.8e-256;			
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						0;
						Gaps
						0;
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Db	1	ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGTCTGATGTGCTG 60	
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Qy	181	GTTGAGGCTCGGAAGGCACTTCTGTGTCAGTTCAAGGCGAGTGTACTACACCAAC 240	
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Qy	241	GGGACGCGCATACGCTCTGTGACAGATACATCTACACCGGAGGAGTACGTGCGC 300	
Db	241	GGGACGCGCATACGCTCTGTGACAGATACATCTACACCGGAGGAGTACGTGCGC 300	
Qy	301	TACGACGCGAGCTGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCAGACGCCGAG 360	
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Qy	481	GTGCGCATCTCCTGTCAGACAGAGCCCTCAACCAACACACACTCTGCTGTGTCG 540	
Db	481	GTGCGCATCTCCTGTCAGACAGAGCCCTCAACCAACACACACTCTGCTGTGTCG 540	
Qy	541	GTGACAGATTTTACCCAGCAAGATCAAGTGGCTGTTTTCAGGAATGGCCAGGAG 600	
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Qy	601	ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGAGCCTTCCAGTCTCTG 660	
Db	601	ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGAGCCTTCCAGTCTCTG 660	
Qy	661	GTATGCTGGAGATGACCCCTCATCAGGAGGCTTACACTGSCCATGTGGAGCATCCC 720	
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Qy	721	AGCCTGAAGAGCCCATCACTGTGAGTGGAGGCAAGTCCGAGTCTGCCCGAGCAAG 780	
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Qy	841	AAGAAAAAGACGCTACGCTGAAGTGAACCTTCAAGCCCTCAAGAAAGAACTGCCCCAG 900	
Db	841	AAGAAAAAGACGCTACGCTGAAGTGAACCTTCAAGCCCTCAAGAAAGAACTGCCCCAG 900	
Qy	901	CATCATCATCATCATCAT 918	
Db	901	CATCATCATCATCATCAT 918	

RESULT 2
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DEFINITION Sequence 25 from patent US 5820866.
ACCESSION AR047947
VERSION AR047947.1 GI:5970290
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

REFERENCE	1 (bases 1 to 893)	Unclassified.
AUTHORS	Kappler, J.W. and Marrack, P.	
TITLE	Product and process for T cell regulation	
JOURNAL	Patent: US 5820866-A 25 13-OCT-1998;	
FEATURES	Location/Qualifiers	
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Best Local Similarity	94.6%; Pred. No. 2.6e-191;	
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Qy	61	AGCAGCCCGGACTAGGCGGAACTCCATCTGCTTTCTGCGGTGCTGAGCACC 120
Db	121	AGCAGCCCGGACTAGGCGGAACTCCATCTGCTTTCTGCGGTGCTGAGCACC 175
Qy	121	ATCGTGTGTCGGCAGCTGGGACGGAGTGGGGCTCACTAGTCCCGAGGCTCTGGA 180
Db	176	ATGAGGCTGGCAG-----AGGAGTGGGGCTCACTAGTCCCGAGGCTCTGGA 225
Qy	181	GTTGAGGCTCGGAAGGCACTTCTGTGTCAGTTCAAGGCGAGTGTACTACACCAAC 240
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Qy	241	GGGACGCGCATACGCTCTGTGACAGATACATCTACACCGGAGGAGTACGTGCGC 300
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Qy	301	TACGACGCGAGCTGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCAGACGCCGAG 360
Db	346	TACGACGCGAGCTGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCAGACGCCGAG 405
Qy	361	TACTGGAACAGCCGCGAGATCTGTGAGCGAACCGCGCGAGTGGACACGCGCTGC 420
Db	406	TACTGGAACAGCCGCGAGATCTGTGAGCGAACCGCGCGAGTGGACACGCGCTGC 465
Qy	421	AGACACAACTACGAGGGCGGAGACAGCACCTCTCCGCGGCTTTGAACAGCCCAAT 480
Db	466	AGACACAACTACGAGGGCGGAGACAGCACCTCTCCGCGGCTTTGAACAGCCCAAT 525
Qy	481	GTGCGCATCTCCTGTCAGGACAGGCCCCTCAACCAACACACTCTGCTGTGTCG 540
Db	526	GTGCGCATCTCCTGTCAGGACAGGCCCCTCAACCAACACACTCTGCTGTGTCG 585
Qy	541	GTGACAGATTTTACCCAGCAAGATCAAGTGGCTGTTTTCAGGAATGGCCAGGAG 600
Db	586	GTGACAGATTTTACCCAGCAAGATCAAGTGGCTGTTTTCAGGAATGGCCAGGAG 645
Qy	601	ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGAGCCTTCCAGTCTCTG 660
Db	646	ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGAGCCTTCCAGTCTCTG 705
Qy	661	GTATGCTGGAGATGACCCCTCATCAGGAGGCTTACACTTGCATGTGGAGCATCCC 720
Db	706	GTATGCTGGAGATGACCCCTCATCAGGAGGCTTACACTTGCATGTGGAGCATCCC 765
Qy	721	AGCCTGAAGAGCCCATCATCTGTGAGTGGAGGCAAGTCCGAGTCTGCCCGAGCAAG 780
Db	766	AGCCTGAAGAGCCCATCATCTGTGAGTGGAGGCAAGTCCGAGTCTGCCCGAGCAAG 825
Qy	781	GGAGTGGAGG 791
Db	826	TAAGCATGCGG 836
RESULT	3	

Db	780	AGCCTGAAGAGCCCATCACTGTGGAGTGGAGGCGACAGTCTGAGTCTGCCCTGGAGCAAG	8339
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DEFINITION	Sequence 123 from patent US 5869270.	linear	PAT 29-SEP-1999
ACCESSION	AR033964		
VERSION	AR033964.1	GI:5949569	
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unknown.		
REFERENCE	Unclassified.		
AUTHORS	1 (bases 1 to 1382)		
TITLE	Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.		
JOURNAL	Single chain MHC complexes and uses thereof		
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Qy	121	ATCGTGTGTCTCGGCGAGCTGGGACGGAGTGGGGGCTCACTAGTGCCTCCGAGGCTCTGGA	180
Db	105	GCTGCTCACGCTGAAATCAACGAAGCTGTGTGCTAGCGAGGGGGCGGAAGCGCGGA	164
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Qy	421	AGACACAATCTACAGGGGGCGGAGACAGCACCTCTCTCGCGCGCTTGAACAGCCCAAT	480
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DEFINITION	Sequence 123 from patent US 6309645.		linear
ACCESSION	ARI75097		
VERSION	ARI75097.1	GI:17916396	
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unknown.		
REFERENCE	1. (bases 1 to 1382)		
AUTHORS	Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.		
TITLE	MHC molecules and uses thereof		
JOURNAL	Patent: US 6309645-A 123 30-OCT-2001;		
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Db	66	AGCAGCCCAAGGAC-----CTTAAGTATCTCTCAGGCTGTTTCC	104
Qy	121	ATCGTGTGTGTCGGCAGCTGGGCGAGGTGGGGGCTCACTAGTGGCCCCGAGGCTCTGGA	180
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Db	225	GGGACGACGGCATACGGCTCTGTACACAGATACATCTACAAACGGGAGGAGTACGTGGC	284
Qy	301	TACACACGACGCTGGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGCCAGAGCCCGAG	360
Db	285	TACACACGACGCTGGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGCCAGACGCCGAG	344
Qy	361	TACTGGAAACAGCCGAGAGTCTGTGAGCGAAACGGGGCGGAGGTGGACACCGCGCTGC	420
Db	345	TACTGGAAACAGCCGAGATCTGTGAGCGAAACGGGGCGGAGGTGGACACCGCGCTGC	404
Qy	421	AGACACAACCTACGAGGGCGGAGACAGACACCTTCCCTGCGCGCGCTTGAACACGCCCAAT	480
Db	405	AGACACAACCTACGAGGGGCGGAGACAGACACCTTCCCTGCGCGCGCTTGAACACGCCCAAT	464
Qy	481	GTCCGCATCTCCCTGTCCAGGACAGAGCCCTCAACCAACACACACTCTGCTGTCTCG	540
Db	465	GTCCGCATCTCCCTGTCCAGGACAGAGCCCTCAACCAACACACACTCTGCTGTCTCG	524
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Db	525	GTGACAGATTTCTACCCAGCCCAAGATCAAGTGGCTGTTTCAGGAATGGCCAGAGAG	584
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Db	585	ACAGTGGGGTCTCATCCACACAGCTATTAGAAATGGGGACGTGGACCTTCCAGGTCCTG	644
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Db	645	GTCAATGCTGGAGATGACCCCTCATACAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC	704
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Db	705	AGCTGAAGAGCCCCCATCACTGTGGAGTGA	735
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VERSION	CS079301.1	GI:63093743	
KEYWORDS	unidentified		
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REFERENCE	1		
AUTHORS	Rhode, P.R., Jiao, J.A., Burkhardt, M. and Wong, H.C.		
TITLE	MHC complexes and uses thereof		
JOURNAL	Patent: EP 1526141-A 123 27-APR-2005;		
FEATURES	Altor Bioscience Corporation (US)		
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Qy	181	GGTGGAGGCTCCAAAGGCATTTCTGTGTCAGTTTCAAGGGCGAGTGCTACTACACCAAC	240
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DEFINITION Sequence 121 from patent US 6309645.
ACCESSION AR175095
VERSION AR175095.1 GI:17916394
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 1385)
Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
TITLE MHC molecules and uses thereof

JOURNAL Patent: US 6309645-A 121 30-OCT-2001;
LOCATION/Qualifiers

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Best Local Similarity 89.6%; Pred. No. 1.7e-165;
Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;
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CS079299 1385 bp DNA linear PAT 06-MAY-2005
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DEFINITION CS079299
ACCESSION CS079299
VERSION CS079299.1 GI:63093741
KEYWORDS
SOURCE unidentified
ORGANISM unclassified sequences.

REFERENCE 1
Rhode,P.R., Jiao,J.A., Burkhardt,M. and Wong,H.C.
AUTHORS MHC complexes and uses thereof
TITLE Patent: EP 1526141-A 121 27-APR-2005;
JOURNAL Altor Bioscience Corporation (US)
LOCATION/Qualifiers

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Db	585	ACAGTGGGGTCTCATCCACACAGCTTTATTAGGAATGGGACTGGACCTTCCAGGTCCTG		644
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Db	645	GTCACTGTGGAGATGACCCCTCATCAGGAGAGGTCTACACTGCGCATGTGGAGCATCCC		704
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ARI52030

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Sequence 24 from patent US 6232445.

ARI52030.1

GI:151118080

Unknown.

Unknown.

Unclassified.

1 (bases 1 to 1508)

Rhode, P.R., Acevedo, J., Burkhardt, M., Jiao, J.-a. and Wong, H.C.

Soluble MHC complexes and methods of use thereof

Patent: US 6232445-A 24 15-MAY-2001;

Location/Qualifiers

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linear

PAT 08-AUG-2001

Query Match

Best Local Similarity

Matches

673; Conservative

66.2%; Score 607.8; DB 2; Length 1508;

89.6%; Pred. No. 1.8e-165;

0; Mismatches 57; Indels 21; Gaps 1

ORIGIN

source

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Query Match      66.2%; Score 607.8; DB 2; Length 1508;
Best Local Similarity 89.6%; Pred. No. 1.8e-165;
Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;

QY 1 ATGGCTCTGAGATCCCGAGCTCTCTCTAGCTGCTGTGGTGGTGTGATGGTCTG 60
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ACCESSION CS079300
VERSION    CS079300.1 GI:63093742
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SOURCE     unidentified
ORGANISM   unclassified sequences.
REFERENCE  1
AUTHORS    Rhode, P.R., Jiao, J.A., Burkhardt, M. and Wong, H.C.
TITLE      MHC complexes and uses thereof
JOURNAL    Patent: EP 1526141-A 122 27-APR-2005;
           Altor BioScience Corporation (US)
FEATURES   Location/Qualifiers

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GenCore version 5.1.9
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8	607.8	66.2	1385	2 AAT86987	Aat86987 SSC1 sing
9	607.8	66.2	1385	8 ACA60742	ACA60742 Mouse MHC
10	607.8	66.2	1508	2 AAT86988	Aat86988 SCTL sing
11	607.8	66.2	1508	2 AAX89069	Aax89069 Single ch
12	607.8	66.2	1508	8 ACA60743	ACA60743 Mouse MHC
13	606.2	66.0	1382	2 AAT17588	Aat17588 Vector SC
14	606.2	66.0	1385	2 AAT17586	Aat17586 Vector SS
15	606.2	66.0	1508	2 AAT17587	Aat17587 Vector SC
16	598.6	65.2	4724	2 AAV12068	Aav12068 Murine IA
17	561.8	61.2	798	2 ADJ75986	Adj75986 Marker ge
18	561.8	61.2	798	12 ADX26090	Adx26090 Novel cel

19	549.8	59.9	1085	4	ABI99040	Abi99040 Murine pC
20	542	59.0	702	2	AAQ03170	Aaq03170 Sequence
21	542	59.0	702	2	AAT06286	Aat06286 I-Ab-beta
22	542	59.0	702	2	AAQ56920	Aaq56920 Mouse I-A
23	535.6	58.3	702	2	AAQ35055	Aaq35055 IAB beta
24	525.4	57.2	1698	4	ABI99038	Abi99038 Murine pC
25	502.6	54.7	1243	6	ABN84048	Abn84048 Single ch
26	499.4	54.4	1662	4	ABI99039	Abi99039 Murine pC
27	497.2	54.2	1686	4	ABI99031	Abi99031 MBP 1-14
28	497.2	54.2	1701	4	ABI99028	Abi99028 IAS MBP 1
29	497.2	54.2	2059	4	ABI99032	Abi99032 MBP 1-14
30	497.2	54.2	2346	4	ABI99027	Abi99027 IAS MBP 1
31	486	54.0	1680	4	ABI99021	Abi99021 I-As MBP
32	496	54.0	1707	4	ABI99030	Abi99030 IAS MBP 9
33	496	54.0	2053	4	ABI99029	Abi99029 IAS MBP 9
34	496	54.0	2343	4	ABI99033	Abi99033 MBP 90-10
35	433.6	47.2	562	6	ABK63510	Abk63510 Rat seque
36	433.6	47.2	562	10	ADB57995	Adb57995 Toxicity-
37	433.6	47.2	562	10	ABT411775	Abt411775 Toxicity
38	433.6	47.2	562	11	ADM21868	Adm21868 Rat hepat
39	433.6	47.2	562	13	ADV40851	Adv40851 Rat cardi
40	433.6	47.2	562	14	ADX25826	Adx25826 Novel cel
41	414.8	45.2	1869	13	ADQ38634	Adq38634 Human SNP
42	414.8	45.2	1892	13	ADQ38637	Adq38637 Human SNP
43	410.8	44.7	1171	6	ABK84087	Abk84087 Human cDN
44	410.8	44.7	1199	8	ABX63009	Abx63009 Human cDN
45	407.6	44.4	1192	10	AAD63150	Aad63150 Human maj

ALIGNMENTS

RESULT 1
AAAF5099
ID AAF55099 standard; DNA; 921 BP.

AC AAF55099;

DT 15-MAY-2001 (first entry)

DE DNA encoding a fusion protein comprising a beta chain of MHC.

KW Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
KW major histocompatibility complex; Fc region; antigen; T lymphocyte;
KW immunostimulant; vaccine; infection; tumour; ss.

OS Synthetic.

FX Key Location/Qualifiers
CDS 1..921
FT /*tag= a

PN WO200109194-A1.

XX 08-FEB-2001.

PD 28-JUL-2000; 2000WO-FR002193.

XX 29-JUL-1999; 99FR-00009862.

XX (CNRS) CNRS CENT NAT RECH SCI.

PA Glaichenhaus N, Malherbe L;

XX WPI; 2001-182944/18.

DR P-PSDB; AAB67481.

PT New soluble recombinant protein, useful e.g. as immunostimulant,
PT comprises dimeric major histocompatibility complex molecule fused to
PT immunoglobulin Fc region.

XX Example 1; Page 34-35; 43pp; French.

CC The specification describes soluble recombinant proteins that comprise at
CC least a dimer formed from the alpha and beta-chains of MHC (major
CC histocompatibility complex) Class I and II molecules in which at least
CC one chain has, attached to its C-terminus, at least part of the Fc region
CC of an immunoglobulin. The recombinant proteins, when linked to an
CC antigenic peptide, are used to count and/or purify antigen-reactive T
CC lymphocytes and to characterize their phenotype, e.g. in preclinical
CC evaluation of vaccines. They are also used as immunostimulants,
CC particularly for vaccine development (against infections and tumours), to
CC count and determine phenotype of autoreactive T cells in subjects with,
CC or at risk of developing, autoimmune diseases, e.g. for staging or
CC evaluating treatments, and (to purify and/or enrich Ag-reactive T cells
CC from cell cultures or patient samples, for use in subsequent curative or
CC preventative cellular therapy. The present sequence encodes a recombinant
CC protein of the invention, comprising a beta chain of MHC molecules
XX

Sequence 921 BP; 214 A; 265 C; 286 G; 156 T; 0 U; 0 Other;

Query Match 100.0%; Score 918; DB 5; Length 921;
Best Local Similarity 100.0%; Pred. No. 7.3e-203; Mismatches 0; Gaps 0;
Matches 918; Conservative 0; Indels 0;

Qy 1 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGTGCTG 60
Db 1 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGTGCTG 60
Qy 61 AGCAGCCCGGAGCTGAGGGGGAACCTCATCTCTCTGCTCTGCTGCTGAGCACC 120
Db 61 AGCAGCCCGGAGCTGAGGGGGAACCTCATCTCTCTGCTCTGCTGCTGAGCACC 120
Qy 121 ATCTGTGTCTCGGAGCTGGGAGCGAGTGGGGCTCACTAGTGCCTCCGAGGCTCTG 180
Db 121 ATCTGTGTCTCGGAGCTGGGAGCGAGTGGGGCTCACTAGTGCCTCCGAGGCTCTG 180
Qy 181 GGTGAGGCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTCTACTACACCA 240
Db 181 GGTGAGGCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTCTACTACACCA 240
Qy 241 GGGAGCGAGCGATACGGCTGTGACCAAGTACATCTCAACCCGGAGGAGTACGTGG 300
Db 241 GGGAGCGAGCGATACGGCTGTGACCAAGTACATCTCAACCCGGAGGAGTACGTGG 300
Qy 301 TACGACGAGCTGGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCAGACGCCGAG 360
Db 301 TACGACGAGCTGGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCAGACGCCGAG 360
Qy 361 TACTGGAAACAGCCAGCGAGATCTGGAGCGAAGCGCGGCGGAGGTGGACACGCG 420
Db 361 TACTGGAAACAGCCAGCGAGATCTGGAGCGAAGCGCGGCGGAGGTGGACACGCG 420
Qy 421 AGACACACTAGGAGGCGGAGACACGACCTCTCCCTGCGGCGCTTGAACAGCCCA 480
Db 421 AGACACACTAGGAGGCGGAGACACGACCTCTCCCTGCGGCGCTTGAACAGCCCA 480
Qy 481 GTGCGCATCTCCCTGTCAGACAGAGCGCTTCAACACCAACACTCTGCTGTGTCG 540
Db 481 GTGCGCATCTCCCTGTCAGACAGAGCGCTTCAACACCAACACTCTGCTGTGTCG 540
Qy 541 GTGACAGATTTTACCCAGCAAGATCAAAAGTGGCGCTGTTCAAGAAATGGCCAG 600
Db 541 GTGACAGATTTTACCCAGCAAGATCAAAAGTGGCGCTGTTCAAGAAATGGCCAG 600
Qy 601 ACAGTGGGGGTCTCATCCACAGCTTATAGGAATGGGACTGGACCTTCCAGGTCT 660
Db 601 ACAGTGGGGGTCTCATCCACAGCTTATAGGAATGGGACTGGACCTTCCAGGTCT 660
Qy 661 GTCATGTCTGGAGATGACCCCTCATCAGGAGAGGTCTACACTGCTGTGGAGCATCC 720
Db 661 GTCATGTCTGGAGATGACCCCTCATCAGGAGAGGTCTACACTGCTGTGGAGCATCC 720
Qy 721 AGCTTGAAGAGCCCCATCACTGTGAGTGGAGGGCACAGTCCGAGTCTGCCCGGAG 780
Db 721 AGCTTGAAGAGCCCCATCACTGTGAGTGGAGGGCACAGTCCGAGTCTGCCCGGAG 780

Qy 781 GGAGGTGGAGGATCCACTACAGTCCCATCAGCTCAGTTGAAAAAGAAATTCGAAGCACTG 840
Db 781 GGAGGTGGAGGATCCACTACAGTCCCATCAGCTCAGTTGAAAAAGAAATTCGAAGCACTG 840
Qy 841 AAGAAAAAGAACGCTCAGCTCAAGTGGAAACTTCAAGGCCCTCAAGAGAAACTCCGCCAG 900
Db 841 AAGAAAAAGAACGCTCAGCTCAAGTGGAAACTTCAAGGCCCTCAAGAGAAACTCCGCCAG 900
Qy 901 CATCATCATCATCATCAT 918
Db 901 CATCATCATCATCATCAT 918
RESULT 2
AAT04262
ID AAT04262 standard; DNA; 893 BP.
XX
AC AAT04262;
DT 16-APR-1996 (first entry)
XX
DE Hybrid IA beta chain gene.
XX
KW Polymerase chain reaction; PCR; primer; amplify;
KW major histocompatibility complex; MHC; T-cell receptor; TCR;
KW autoimmune disease; immunodeficiency disease; immune response;
XX immunoproliferation disease; graft-host rejection; therapy; ss.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT primer_bind 1..16
FT /tag= a
FT /note= "probable primer binding site (primer #233)"
FT primer_bind complement(45..74)
FT /tag= b
FT /note= "binding site for primer #261 (see AAT04260)"
FT CDS 61..828
FT /tag= c
FT /product= "hybrid IA beta chain"
FT sig_peptide 61..141
FT /tag= d
FT /note= "leader region"
FT primer_bind complement(119..172)
FT /tag= e
FT /note= "binding site for primer #331 (see AAT04261)"
FT primer_bind complement(158..212)
FT /tag= f
FT /note= "primer #332 binding site"
FT primer_bind complement(199..250)
FT /tag= g
FT /note= "primer #333 (see AAT04263) binding site"
FT misc_feature 371..389
FT /tag= h
FT /note= "probable primer binding site (primer #270)"
FT mat_peptide 511..825
FT /tag= i
FT /product= "IA beta chain beta 2 region"
FT primer_bind 521..550
FT /tag= j
FT /note= "probable primer binding site (primer #271)"
FT primer_bind 532..554
FT /tag= k
FT /note= "probable primer binding site (primer #272)"
FT primer_bind 808..836
FT /tag= l
FT /note= "probable primer binding site (primer #259)"
FT primer_bind 877..893
FT /tag= m
FT /note= "probable primer binding site (primer #232)"
XX
PN W09523814-A1.

CC acid sequence comprising one or more deletions, substitutions or
CC additions. The molecule of the invention may be useful for detecting an
CC antigen-specific CD4+ T-cell by flow cytometry and for presenting a
CC microorganism-derived mucous membrane invasive protein as an antigen. The
CC method of the invention enables efficient detection of antigen-specific
CC activation of CD4+ T-cells in the mucous membrane. The current sequence
CC is that of the class II major histocompatibility complex-related I-
CC Ab(alpha)-Cholera toxin B subunit (CTB)-leucine zipper (LZ)-BirA fusion
CC cDNA of the invention.
XX
SQ Sequence 945 BP; 230 A; 256 C; 294 G; 165 T; 0 U; 0 Other;

Query Match 72.5%; Score 665.2; DB 12; Length 945;
Best Local Similarity 86.6%; Pred. No. 3e-144;
Matches 759; Conservative 0; Mismatches 108; Indels 9; Gaps 2;
QY 43 GTGGTCTGATGGTCTGAGCAGCCCGGGACTGAGGGCGGAAATCCATCTGCTTCTCG 102
DB 49 GTGACACTGATGGTCTGAGCTCCCACTGGCTTTGGCTGGAGACTCTGCGTGGGAAC 108
QY 103 CGGTCTGAGACACCCGATCGTGTGTCGCGCAGCTGGAGCGAGGTGGGGGCTCACTA 162
DB 109 AATAAGACCGCGCAGCGCATCGCGCCATCAGCATGGCGAAACGAGGTGGTGGTCCGGT 168
QY 163 GTGCCCCGAGGCTCTGGAGGTGGAGCTCCGAAAGGCATTTCGTGTCAGTTCAAGGCG 222
DB 169 GGAGGGGAAG---TGGAGGTGGAGGGTCTGAAAGGCATTTCGTGTACAGTTCAATGGCG 225
QY 223 GAGTCTCTACTACACCAACGGGACGCGGCATACCGCTCGTGACCAGATACATCTACAAAC 282
DB 226 GAGTCTACTTACCAACGGGACGCGGCATACGATATGTGACCAGATACATCTACAAAC 285
QY 283 CGGAGGAGTACGTGGCTTACAGACGACAGTGGGCGAGTACCGCGCGGTGACCGAGCTG 342
DB 286 CGGAGGAGTACGTGGCTTACAGACGACAGTGGGCGAGCACCGCGCGGTGACCGAGCTG 345
QY 343 GGGCGGCGAGCGCGGAGTCTGGAACAGCCAGCGAGATCTCGGAGCAACCGGCGG 402
DB 346 GGGCGGCGAGCGCGGAGTCTGGAACAGCCAGCGGAGATCTCGGAGCAACCGGCGG 405
QY 403 GAGGTGGACACCGCGTGCAGACACAACTACGAGGGGCGGAGCACGACCTCCCTGCGG 462
DB 406 GAGCTGGACACCGGTGTCAGACACAACTACGAGGGGCGGAGCACCACTCCCTGCGG 465
QY 463 CGGCTTGAAACAGCCCAATGTGCCATCTCCCTGTCCAGACAGAGGCGCTCAACACAC 522
DB 466 CGGCTTGAAACAGCCCAATGTGCTATCTCCCTGTCCAGACAGAGGCGCTCAACACAC 525
QY 523 AACCTCTGGTCTGTTCCGTGACAGATTCTACCCAGCCCAAGATCAAAAGTGGCTGTTTC 582
DB 526 AACCTCTGGTCTGTTCCGTGACAGATTCTTACCCAGCCCAAGATCAAAAGTGGCTGTTTC 585
QY 583 AGGAAATGGCCAGGAGACAGTGGGGGTCTCATCCACACAGCTTTATTAGGAATGGGAC 642
DB 586 CGGAATGGCCAGGAGACAGTGGGGGTCTCATCCACACAGCTTTATTAGGAATGGGAC 645
QY 643 TGGACCTTCCAGGCTCTGTATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACC 702
DB 646 TGGACCTTCCAGGCTCTGTATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACC 705
QY 703 TGCCATGTGGAGCATCCAGCTGAAGAGCCCATCACTGTGGAGTGGAGGCGCAGTCC 762
DB 706 TGTACGTGGAGCATCCAGCTGAAGAGCCCATCACTGTGGAGTGGAGGCGCAGTCC 765
QY 763 GAGTCTGCGCGAGCAAGGAGTGGAGGATCCACTACAGCTTCCATCAGCTCAGTTGAAA 822
DB 766 TCAGCAGACC-----TGGTTCGCGCGGATCCACTACAGCTTCCATCAGCTCAGTTGAAA 819
QY 823 AAGAAATTCAGACACTGAGAAAAAAGAACGCTCAGCTGAAGTGAAGAACTTCAAGCCCTC 882
DB 820 AAGAAACTGCAGGCACTTAAAGAAAAAGAACGCTCAGCTGAAGTGAAGAACTTCAAGCCCTC 879
QY 883 AAGAAGAAACTCGCCGAGCATCATCATCATCATCATCATCATCATCATCATCATCATCAT 918

DB 880 AAGAAGAAACTCGCCAGCTGCATCATCATCATCATCATCATCATCATCATCATCATCAT 915
RESULT 4
AAT04269
ID AAT04269 standard; DNA; 1013 BP.
XX
AC AAT04269;
XX
DT 16-APR-1996 (first entry)
XX
DE Hybrid IA beta chain gene.
XX
KW Major histocompatibility complex; MHC; T-cell receptor; TCR;
KW autoimmunity disease; immunodeficiency disease; immune response;
KW immunoproliferation disease; graft-host rejection; therapy; B cell;
KW M12.C3; pM12-IAB-Ea; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT primer_bind 1..18
FT /tag= a
FT /note= "probable primer binding site (primer #76)"
FT primer_bind complement(40..74)
FT /tag= b
FT /note= "binding site for primer #362 (see AAT04270)"
FT CDS 63..959
FT /tag= c
FT /product= "hybrid IA beta chain"
FT sig_peptide 63..143
FT /tag= d
FT /note= "leader region"
FT primer_bind complement(140..191)
FT /tag= e
FT /note= "binding site for primer #363 (see AAT04271)"
FT primer_bind complement(177..226)
FT /tag= f
FT /note= "primer #364 binding site"
FT primer_bind complement(212..266)
FT /tag= g
FT /note= "primer #365 (see AAT04272) binding site"
FT primer_bind 385..403
FT /tag= h
FT /note= "probable primer binding site (primer #270)"
FT mat_peptide 531..959
FT /tag= i
FT /product= "IA beta chain beta 2 region"
FT primer_bind 535..564
FT /tag= j
FT /note= "probable primer binding site (primer #271)"
FT primer_bind 544..568
FT /tag= k
FT /note= "probable primer binding site (primer #272)"
FT primer_bind 823..850
FT /tag= l
FT /note= "probable primer binding site (primer #259)"
FT primer_bind 942..976
FT /tag= m
FT /note= "probable primer binding site (primer #366)"
FT primer_bind 1000..1013
FT /tag= n
FT /note= "probable primer binding site (primer #59)"
XX
PN WO9523814-A1.
XX
PD 08-SEP-1995.
XX
PF 03-MAR-1995; 95WO-US002689.
XX
PR 04-MAR-1994; 94US-00207481.
XX

Best Local Similarity 91.9%; Pred. No. 8.9e-136; Matches 677; Conservative 0; Mismatches 54; Indels 6; Gaps 1;

Qy 182 GTGAGGCTCCGAAAGGCAATTCGTGGTCCAGTTTCAAGGGGAGTGCTACTACACCAACG 241

Db 155 GAGGTGGTAGTGAAAGGCAATTCGTGTACCAAGTTTCATGGCGAGTGTCTACTTACCAACG 214

Qy 242 GGACGACGGCATACGGCTCGTGACCATATCTACAACGGGAGGAGTACGTGGCT 301

Db 215 GGACGACGGCATACGATATGTGACCATATCTACAACGGGAGGAGTACGTGGCT 274

Qy 302 ACGACAGCGCTGGGCGGAGTACCGCGGTGACCGAGCTGGGGCGGCACACGCCAGT 361

Db 275 ACGACAGCGCTGGGCGGAGACCGCGGTGACCGAGCTGGGGCGGCACACGCCAGT 334

Qy 362 ACTGGAACAGCCAGCGGAGATCTCTGGAGCGAAACGCGGGCCGAGGTGGACACGGGTGCA 421

Db 335 ACTGGAACAGCCAGCGGAGATCTCTGGAGCGAAACGCGGGCCGAGGTGGACACGGGTGCA 394

Qy 422 GACACAACCTACAGGGGCGGAGACGACACTCTCCCTGGGGGGCTTGAACAGGCCAATG 481

Db 395 GACACAACCTACAGGGGCGGAGACCCACACTCCCTGGGGGGCTTGAACAGGCCAATG 454

Qy 482 TCGCCATCTCCCTGTCCAGGACAGAGGCCCTCAACCCACACAACCTCTGTGTCTGG 541

Db 455 TCGTCATCTCCCTGTCCAGGACAGAGGCCCTCAACCCACACAACCTCTGTGTCTGG 514

Qy 542 TGACAGATTTCTACCCAGCCAAAGATCAAAAGTGCCTGTTTCAAGGAATGCCAGGAGAGA 601

Db 515 TGACAGATTTCTACCCAGCCAAAGATCAAAAGTGCCTGTTTCCGGAATGCCAGGAGAGA 574

Qy 602 CAGTGGGGTCTCATCCACAGCTTATTAGGAATGGGAGCTTGACCTTCCAGTCTCTGG 661

Db 575 CGGTGGGGTCTCATCCACAGCTTATTAGGAATGGGAGCTTGGACCTTCCAGTCTCTGG 634

Qy 662 TCATGCTCGAGATGACCCCTCATCAGGAGAGGTTCTACACCTGCCATGTGGAGCATCCCA 721

Db 635 TCATGCTCGAGATGACCCCTCTGGGAGTGGAGGAGGTTCTACACCTGTCACTGGAGCATCCCA 694

Qy 722 GCCTGAAGAGCCCATCATCTGTGGAGTGGAGGACAGTCCGAGTCTGCCCGAGCAAG 781

Db 695 GCCTGAAGAGCCCATCATCTGTGGAGTGGAGGACAGTCTGTCAGCAGACC-----TGG 748

Qy 782 GAGGTGGAGGATCCACTACAGTCCATCAGCTCAGTTGAAAAAGAAATTCAGCAGCTGA 841

Db 749 TTCCGCGGGATCCACTACAGTCCATCAGCTCAGTTGAAAAAGAAATTCAGCAGCTGA 808

Qy 842 AGAAAAAGAACGCTCAGCTGAAGTGGAACCTTCAAGCCCTCAAGAAAGAACTCGCCCCAG 901

Db 809 AGAAAAAGAACGCTCAGCTGAAGTGGAACCTTCAAGCCCTCAAGAAAGAACTCGCCCCAG 868

Qy 902 ATCATCATCATCATCAT 918

Db 869 TGCATCATATCTGGAT 885

RESULT 6

AAT86989

ID AAT86989 standard; DNA; 1382 BP.

XX

AC AAT86989;

XX

DT 27-MAR-1998 (first entry)

XX

DE SCE1 single chain gene.

XX

KW Construction; major histocompatibility complex; MHC; fusion complex;

KW SCE1 single chain gene; ss.

XX

OS Synthetic.

XX

FH Key

FT CDS

Location/Qualifiers

6..1382

FT

XX

FN MO9728191-A1.

XX

PD 07-AUG-1997.

XX

PF 30-JAN-1997; 97WO-US001617.

XX

PR 31-JAN-1996; 96US-00596387.

XX

PA (DADE-) DADE INT INC.

XX

XX Rhode PR, Jiao J, Burkhardt M, Wong HC;

PI WPI; 1997-402555/37.

XX

DR P-PSDB; AAW29214.

XX

PT Single chain major histocompatibility complex comprising linked alpha and beta chains - useful for suppressing an immune response to an auto:immune disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes mellitus, etc.

PT

PT

XX

PS Example 17; Page 140-141; 217pp; English.

XX

CC The present sequence was used in the construction of major histocompatibility complex (MHC) fusion complexes

CC

XX

SQ Sequence 1382 BP; 320 A; 373 C; 405 G; 284 T; 0 U; 0 Other;

Query Match 66.2%; Score 607.8; DB 2; Length 1382;

Best Local Similarity 89.6%; Pred. No. 6.5e-131;

Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;

Qy 1 ATGGCTCTGCAGATCCCCAGCTCTCTCTCAGCTGCTGTGTGTGTCTGATGTGTCTG 60

Db 6 ATGGCTCTGCAGATCCCCAGCTCTCTCTCAGCTGCTGTGTGTGTCTGATGTGTCTG 65

Qy 61 AGCAGCCCGGAGTGGGGGGAACCTCCATCTGCTTCTCGCGCTCGCTGAGCACCCG 120

Db 66 AGCAGCCCAAGGAC-----CTTAAATATCTCTCAGCTGTTCAC 104

Qy 121 ATCGTGTGTCCGCGAGCTGGGACGAGGTGGGGCTCACTAGTGCCCGAGGCTCTGGA 180

Db 105 GCTCTCAGCTGAATCAACGAAGTGTGTGTCTAGCGAGGGGGCGGAAGCGCGGA 164

Qy 181 GGTGGAGGCTCCGAAAGGCATTTCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 240

Db 165 GGGGGAACCTCCGAAAGGCATTTCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 224

Qy 241 GGGACGAGCGCATACCGCTCTGCACCATGATATCTACAAACCGGAGGAGTACGTGCGC 300

Db 225 GGGACGAGCGCATACCGCTCTGCACCATGATATCTACAAACCGGAGGAGTACGTGCGC 284

Qy 301 TACGACAGCGAGTGGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGGCAGACCCGAG 360

Db 285 TACGACAGCGAGTGGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGGCAGACCCGAG 344

Qy 361 TACTGGAAACGCCAGCGGAGATCTCTGAGGGAACCGGGCGGAGTGGGACACCGCGTGC 420

Db 345 TACTGGAAACGCCAGCGGAGATCTCTGAGGGAACCGGGCGGAGTGGGACACCGCGTGC 404

Qy 421 AGACACAACCTACGAGGGGCGGAGACCACTCCCTCGCGCGCTTTGAACAGCCCAAT 480

Db 405 AGACACAACCTACGAGGGGCGGAGACCACTCCCTCGCGCGCTTTGAACAGCCCAAT 464

Qy 481 GTCCGCACTCTCCCTGTCCAGGACAGAGGCCCTCAACCAACCAACACTCTGTGTGTGT 540

Db 465 GTCCGCACTCTCCCTGTCCAGGACAGAGGCCCTCAACCAACCAACACTCTGTGTGTGT 524

Qy 541 GTGACAGATTTCTACCCAGCCCAAGATCAAAAGTGGCTGTTCAGGAATGGCCAGGAGG 600

Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAAAGTGGCTGTTCAGGAATGGCCAGGAGG 584

QY	601	ACAGTGGGGGCTTCATCCACACAGCTTATAGGAATGGGACTTGGACCTTCAGAGTCCCTG	660
Db	595	ACAGTGGGGGCTTCATCCACACAGCTTATAGGAATGGGACTTGGACCTTCAGAGTCCCTG	644
QY	661	GTTCATCTGGAGATGATCCCCCTCATCAGGAGAGGTTCTACCTGCCATGTGAGCATCCC	720
Db	645	GTTCATCTGGAGATGATCCCCCTCATCAGGAGAGGTTCTACCTGCCATGTGAGCATCCC	704
QY	721	AGCCTGAAGAGCCCCCATCACTGTGGAGTGGGA	751
Db	705	AGCCTGAAGAGCCCCCATCACTGTGGAGTGGGA	735

RESULT 7

ACA60744
ID ACA60744 standard; DNA: 1382 BP.

ACA60744:

DT 16-JUN-2003 (first entry)

DE Mouse MHC I-Ad/Ova 323-339 synthetic gene SCEL.

KW MHC; major histocompatibility complex; gene therapy; fusion complex;
 KW peptide-binding groove; T cell modulation; class II MHC; vaccine;
 KW autoimmune disorder; multiple sclerosis; rheumatoid arthritis;
 KW insulin-dependent diabetes mellitus; myasthenia gravis; immunogen;
 KW chronic allergy; mouse; ds; I-Ad; gene.

OS Mu8 8D.

OS	maas sp.
OS	Synthetic.

PN US2002198144-A1.

26-DEC-2002.

06-JUL-2001: 2001US-009000379.

XX
PR 29-JUL-1994: 94US-00283302

23-000-1337, 3#03-00283302, 95US-00382454.
PR 01-FEB-1995: PR

PR 17-JAN-1997; 97US-00776084.

PA (DADE-) DADE INT INC.

PI Wong HC. Rhode PR. Weidanz JA. Grammer S. Edwards AC.

PI Chavallaz P, Jiao JJ:

DR WPI: 2003-341126/32.

DR P-PSDB; ABU72108.

PT Novel major histocompatibility complex fusion complex having presenting
PT peptide covalently linked to MHC molecule containing peptide-binding
PT groove, used for suppressing immune response in multiple sclerosis,
PT allergies.

Example 17: Fig 29: 126pp: English

The invention relates to a major histocompatibility complex (MHC) fusion complex (I) comprising an MHC molecule that contains a peptide-binding groove, and a presenting peptide covalently (e.g. an antigenic peptide) linked to the MHC molecule, where (I) is capable of modulating the activity of a T cell. Also included are a DNA construct coding for the complex, where the MHC molecule is a class II MHC (e.g. mouse I-Ad or I-Es, or human HLA-DR1 (human leukocyte antigen-DR1)), a multivalent MHC fusion complex comprising two or more linked complexes, identifying a peptide that can modulate the activity of T cells (involving introducing into host cells cloning vectors that each contain the fusion complex DNA, culturing the host cells under conditions suitable for expression of the MHC fusion complex, and selecting host cells that express MHC fusion complex that modulate the activity of T cells), a single recombinant expression vector comprising DNA that codes for the alpha and beta chains of the fusion complex MHC protein, a single recombinant expression vector comprising DNA that codes for a T cell costimulatory factor and the alpha

```
Db      645  GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy      721  AGCTGAAGAGCCCCCATCACTGTGGAGTGA 751
Db      705  AGCCTGAAGAGCCCCCATCACTGTGGAGTGA 735

RESULT 8
ID      AAT86987 standard; DNA; 1385 BP.
XX
AC      AAT86987;
XX
DT      27-MAR-1998 (first entry)
XX
DE      SSI1 single chain gene.
XX
KW      Construction; major histocompatibility complex; MHC; fusion complex;
KW      SSI1 single chain gene; ss.
XX
OS      Synthetic.
XX
FH      Key      Location/Qualifiers
FT      CDS      6..1385
FT      FT      /*tag= a
XX
PN      WO9728191-A1.
XX
PD      07-AUG-1997.
XX
PF      30-JAN-1997; 97WO-US001617.
XX
PR      31-JAN-1996; 96US-00596387.
XX
PA      (DADE-) DADE INT INC.
XX
PI      Rhode PR, Jiao J, Burkhardt M, Wong HC;
XX
DR      WPI; 1997-402555/37.
XX
P-PSDB; AAW29212.
XX
Single chain major histocompatibility complex comprising linked alpha and
PT beta chains - useful for suppressing an immune response to an auto:immune
PT disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes
PT mellitus, etc.
XX
PS      Example 17; Page 135-137; 217pp; English.
XX
CC      The present sequence was used in the construction of major
CC      histocompatibility complex (MHC) fusion complexes
XX
SQ      Sequence 1385 BP; 316 A; 383 C; 399 G; 287 T; 0 U; 0 Other;

Query Match      66.2%; Score 607.8; DB 2; Length 1385;
Best Local Similarity 89.6%; Pred No. 6.5e-131;
Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;

Qy      1  ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGGTCTGATGGTCTG 60
Db      6  ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGGTCTGATGGTCTG 65
Qy      61  AGCAGCCCGGAGCTGAGGGGGAACTCCATCTGCTTCTCGCCCTCGCTGGAGCACCG 120
Db      66  AGCAGCCCAAGGAC-----CTTAAGTATCTCTCAGGCTGTTTAC 104
Qy      121  ATCGTGGTGTCCGGCAGCTGGGACGAGGTGGGGCTCACTAGTCCCGGAGGCTCTGGA 180
Db      105  GCTGTCTACGCTGAAATCAACGAACTGCTGTCTAGCGAGGGGGCGGAGCGCGGA 164
Qy      181  GGTGGAGGCTCCGAAAGGCAATTCGTGTGTCAGTTCAAGGGCGAGTGTACTACACCAAC 240
Db      165  GGGGGAAATCCGAAAGGCAATTCGTGTGTCAGTTCAAGGGCGAGTGTACTACACCAAC 224
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Qy      241  GGGAGCGAGCGCATACGGCTCGTGACAGATACATCTACAAACCGGGAGGAGTACGTGGCG 300
Db      225  GGGAGCGAGCGCATACGGCTCGTGACAGATACATCTACAAACCGGGAGGAGTACGTGGCG 284
Qy      301  TACGACAGCAGCTGGGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGGCCAGACGCCAG 360
Db      285  TACGACAGCAGCTGGGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGGCCAGACGCCAG 344
Qy      361  TACTGGAACAGCCAGCCGAGATCTCTGAGGGAACCGGGCGGAGGTGGACACGGCGTGC 420
Db      345  TACTGGAACAGCCAGCCGAGATCTCTGAGGGAACCGGGCGGAGGTGGACACGGCGTGC 404
Qy      421  AGACACAACCTACGAGGGGCGGAGACGACGACCTCCCTGCGGCGGCTTTGAACAGGCCAAT 480
Db      405  AGACACAACCTACGAGGGGCGGAGACGACGACCTCCCTGCGGCGGCTTTGAACAGGCCAAT 464
Qy      481  GTGCCCATCTCCCTGTCCAGGACAGAGGCCCTCAACCAACCAACACACTCTCTGGTCTGTCG 540
Db      465  GTGCCCATCTCCCTGTCCAGGACAGAGGCCCTCAACCAACCAACACACTCTCTGGTCTGTCG 524
Qy      541  GTGACAGATTCTACCCAGCCAGATCAAGTGGCTGGTTTCAGGAAATGGCCAGGAGGAG 600
Db      525  GTGACAGATTCTACCCAGCCAGATCAAGTGGCTGGTTTCAGGAAATGGCCAGGAGGAG 584
Qy      601  ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCCAGGTCTCTG 660
Db      585  ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCCAGGTCTCTG 644
Qy      661  GTCATGCTGGAGATGACCCCTCATCAGGAGAGTCTACACCTGCCATGTGGAGCATCCC 720
Db      645  GTCATGCTGGAGATGACCCCTCATCAGGAGAGTCTACACCTGCCATGTGGAGCATCCC 704
Qy      721  AGCCTGAAGAGCCCATCACTGTGGAGTGA 751
Db      705  AGCCTGAAGAGCCCATCACTGTGGAGTGA 735

RESULT 9
ID      ACA60742 standard; DNA; 1385 BP.
XX
AC      ACA60742;
XX
DT      16-JUN-2003 (first entry)
XX
DE      Mouse MHC I-Ad/Ova 323-339 synthetic gene SSI1.
XX
KW      MHC; major histocompatibility complex; gene therapy; fusion complex;
KW      peptide-binding groove; T cell modulation; class II MHC; vaccine;
KW      autoimmune disorder; multiple sclerosis; rheumatoid arthritis;
KW      insulin-dependent diabetes mellitus; myasthenia gravis; immunogen;
KW      chronic allergy; mouse; ds; I-Ad; gene.
XX
OS      Mus sp.
OS      Synthetic.
XX
PN      US2002198144-A1.
XX
PD      26-DEC-2002.
XX
PF      06-JUL-2001; 2001US-00900379.
XX
PR      29-JUL-1994; 94US-00283302.
PR      01-FEB-1995; 95US-00382454.
PR      17-JAN-1997; 97US-00776084.
XX
PA      (DADE-) DADE INT INC.
XX
PI      Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
PI      Chavalliaz P, Jiao JJ;
XX
DR      WPI; 2003-341126/32.
DR      P-PSDB; ABU72106.
```


CC	histocompatibility complex (MHC) fusion complexes	
XX		
SQ	Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;	
	Query Match 66.2%; Score 607.8; DB 2; Length 1508;	
	Best Local Similarity 89.6%; Pred. No. 6.7e-131;	
	Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;	
Qy	1 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGGTCTG 60	
Db	6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGGTCTG 65	
Qy	61 AGCAGCCCGGAGTACGAGCGGAACTCCATCTCTCTCGCGTCTGCTGAGCAGCCG 120	
Db	66 AGCAGCCCAAGAC-----CTTAAGTATCTCTCAGGCTGTTTCTAC 104	
Qy	121 ATCGTGTGTCCGGCAGCTGGAGGTTGGGGCTCCTAGTGTCCCGAGGCTCTGGA 180	
Db	105 GCTGCTCAGCTGAAATCAACGAGCTGTGTCTAGCGAGGGGGCGAAGCGCGGA 164	
Qy	181 GGTGAGGCTCGAAAGGCATTTTCGTGGTCCAGTTCAAGGGCGAGTGTCTATACACCAAC 240	
Db	165 GGGGAAATCCGAAAGGCATTTTCGTGGTCCAGTTCAAGGGCGAGTGTCTATACACCAAC 224	
Qy	241 GGGACGACGCCATACGGCTCGTACCCAGATACATCTACACCCGGGAGGAGTACGTGGCG 300	
Db	225 GGGACGACGCCATACGGCTCGTACCCAGATACATCTACACCCGGGAGGAGTACGTGGCG 284	
Qy	301 TACCACAGCAGCTGGCGAGTACCGCGGTGTACCGAGTGTGGGCGGCCAGACGCCGAG 360	
Db	285 TACCACAGCAGCTGGCGAGTACCGCGGTGTACCGAGTGTGGGCGGCCAGACGCCGAG 344	
Qy	361 TACTGGAACACCCAGCCGAGATCCTGGAGCGAAACGGGGCCGAGGTGACACACGCCGCTGC 420	
Db	345 TACTGGAACACCCAGCCGAGATCCTGGAGCGAAACGGGGCCGAGGTGACACACGCCGCTGC 404	
Qy	421 AGACACAATACGAGGGCCGAGACACACCTCTCCCTGCGCGGCTTGAACAGCCCAAT 480	
Db	405 AGACACAATACGAGGGCCGAGACACACCTCTCCCTGCGCGGCTTGAACAGCCCAAT 464	
Qy	481 GTCGCCATCTCCTGTCCAGGACAGAGCCCTCAACACCAACCACTCTGTGTCTGTTCG 540	
Db	465 GTCGCCATCTCCTGTCCAGGACAGAGCCCTCAACACCAACCACTCTGTGTCTGTTCG 524	
Qy	541 GTGACAGATTTTACCCAGCAAGATCAAGTGGCTGGTTTCAGAAATGGCCAGAGGAG 600	
Db	525 GTGACAGATTTTACCCAGCAAGATCAAGTGGCTGGTTTCAGAAATGGCCAGAGGAG 584	
Qy	601 ACAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGGACTGGACCTTCCAGGTCTCTG 660	
Db	585 ACAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGGACTGGACCTTCCAGGTCTCTG 644	
Qy	661 GTCATGTGGAGATGACCCCTCATCAGGGAGAGGTCTACCTGCGCATGTGGAGATCCC 720	
Db	645 GTCATGTGGAGATGACCCCTCATCAGGGAGAGGTCTACCTGCGCATGTGGAGATCCC 704	
Qy	721 AGCCTGAGAGCCCATCACTGTGAGTGA 751	
Db	705 AGCCTGAGAGCCCATCACTGTGAGTGA 735	
RESULT 11		
AAx89069		
ID	AAx89069 standard; DNA; 1508 BP.	
XX		
AC	AAx89069;	
XX		
DT	14-SEP-1999 (first entry)	
XX		
DE	Single chain IAd/OVA 323-229 MHC fusion protein encoding DNA.	
XX		
KW	Major histocompatibility complex; MHC; single chain MHC; sc-MHC; Ig;	
KW	peptide binding groove; immunoglobulin; T cell receptor; immune response;	

KW	immune-related disorder; antigenic peptide; fusion protein; ss.	
XX		
OS	Synthetic.	
XX		
PN	W09921572-A1.	
XX		
PD	06-MAY-1999.	
XX		
PF	13-OCT-1998; 98WO-US021520.	
XX		
PR	29-OCT-1997; 97US-00960190.	
XX		
PA	(SUNO-) SUNOL MOLECULAR CORP.	
XX		
PI	Rhode PR, Acevedo J, Burkhardt M, Jiao J, Wong HC;	
XX		
DR	WPI; 1999-418411/35.	
DR	P-PSDB; AAY27111.	
XX		
PT	Single chain major histocompatibility complex class I complexes.	
XX		
PS	Example 1; Fig 1; 148pp; English.	
XX		
CC	The invention relates to new single chain major histocompatibility complex (sc-MHC) class II complexes that comprise a peptide binding groove, and a modified class II beta 2 chain or covalently linked immunoglobulin (Ig) light chain constant (C1) region. The MHC complexes are useful for detection and analysis of peptide ligands, pathogenic T-cells, for functional, cellular and molecular assays. They can be used to identify and isolate T cell receptor and/or MHC agonists and antagonists. They can be used in vivo to compete with pathogenic antigen presenting cells involved in immune-related disorders. They can also be used to raise antibodies and to screen immune cells. It is also use in a method of suppressing an immune response in mammals. The sc-MHC complexes comprising modified class II beta 2 chains and/or Ig-C1 regions are soluble and provide enhanced yield. These MHC complexes also can contain single antigenic peptides readily isolated from expressing cells in significant quantities. The polyspecific MHC complexes also provide a means to detect cells expressing multiple target structures with a single complex. The present sequence represents a DNA encoding a single chain IAd/OVA 323-229 MHC fusion protein	
XX		
SQ	Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;	
	Query Match 66.2%; Score 607.8; DB 2; Length 1508;	
	Best Local Similarity 89.6%; Pred. No. 6.7e-131;	
	Matches 673; Conservative 0; Mismatches 57; Indels 21; Gaps 1;	
Qy	1 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGGTCTG 60	
Db	6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGGTCTG 65	
Qy	61 AGCAGCCCGGAGTACGAGGGGAACTCCATCTGCTTCTCGCGTCTGCTGGAGCACCG 120	
Db	66 AGCAGCCCAAGAC-----CTTAAGTATCTCTCAGGCTGTTTCTAC 104	
Qy	121 ATCGTGTGTCCGGCAGCTGGAGGTTGGGGCTCCTAGTGTCCCGAGGCTCTGGA 180	
Db	105 GCTGCTCAGCTGAAATCAACGAGCTGTGTCTAGCGAGGGGGCGAAGCGCGGA 164	
Qy	181 GGTGAGGCTTCGAAAGGCATTTTCGTGGTCCAGTTCAAGGGCGAGTGTCTATACACCAAC 240	
Db	165 GGGGAAATCCGAAAGGCATTTTCGTGGTCCAGTTCAAGGGCGAGTGTCTATACACCAAC 224	
Qy	241 GGGACGACGCCATACGGCTCGTACCCAGATACATCTACACCCGGGAGGAGTACGTGGCG 300	
Db	225 GGGACGACGCCATACGGCTCGTACCCAGATACATCTACACCCGGGAGGAGTACGTGGCG 284	
Qy	301 TACCACAGCAGCTGGCGAGTACCGCGGTGTACCGAGTGTGGGCGGCCAGACGCCGAG 360	
Db	285 TACCACAGCAGCTGGCGAGTACCGCGGTGTACCGAGTGTGGGCGGCCAGACGCCGAG 344	
Qy	361 TACTGGAACACCCAGCCGAGATCCTGGAGCGAAACGGGGCCGAGGTGACACACGCCGCTGC 420	


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Qy 481 GTCGCCATCTCCCTGTCAGACAGAGGCCCTCAACGACCAACACACTCTGGTCTGTTCG 540
Db 465 GTCGCCATCTCCCTGTCAGACAGAGGCCCTCAACGACCAACACACTCTGGTCTGTTCG 524
Qy 541 GTGACAGATTCTACCCAGCCAGATCAAGTGGCGTCTGTTTCAGGAATGGCCAGGAGGAG 600
Db 525 GTGACAGATTCTACCCAGCCAGATCAAGTGGCGTCTGTTTCAGGAATGGCCAGGAGGAG 584
Qy 601 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCCAGTCTCTG 660
Db 585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCCAGTCTCTG 644
Qy 661 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCGCATGTGGAGCATCCC 720
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCGCATGTGGAGCATCCC 704
Qy 721 AGCCTGAAGAGCCCATCACTGTGGAGTGA 751
Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGA 735

RESULT 13
ID AAT17588 standard; DNA; 1382 BP.
XX
AC AAT17588;
XX
DT 26-SEP-1996 (first entry)
XX
DE Vector SCE1-derived single chain gene encoding MHC fusion complex.
XX
KW MHC; major histocompatibility complex; PCR; polymerase chain reaction;
KW T cell activity modulator; antagonist; immune disorder; allergy;
KW multiple sclerosis; insulin-dependent diabetes mellitus;
KW rheumatoid arthritis; myasthenia gravis; ds.
XX
OS Synthetic.
XX
FH Key
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FT 6..86
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FT /note= "murine MHC class II I-Ad gene beta chain leader
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PF 31-JUL-1995; 95WO-US009816.
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PR 29-JUL-1994; 94US-00283302.
XX
PR 01-FEB-1995; 95US-00382454.
XX
PA (DADE-) DADE INT INC.
XX
PI Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
PI Chavallaz P, Jiao J;
XX
XX WPI; 1996-129343/13.
XX P-PSDB; AAR98907.
XX
XX Major histocompatibility complex fusion complex for modulating T cell
XX activity - used in the treatment of immune disorders, e.g. multiple
XX sclerosis, IDDM and rheumatoid arthritis.
XX
XX Example 17; Fig 29; 210pp; English.
XX
XX AAT17588 encodes a murine MHC fusion complex capable of modulating T cell
XX activity encoded by the vector SCE1. The MHC fusion complex comprises at
XX least one MHC molecule containing a peptide-binding groove and a
XX presenting peptide covalently linked to the MHC molecule and opt. a
XX transmembrane domain. DNA encoding a MHC fusion complex may be cloned
XX into a host cell to express the complex. The transformed cells may then
XX be used to identify peptides that modulate, pref. antagonise, T cell
XX activity. DNA encoding a MHC fusion complex or a single chain fusion
XX molecule may be used to vaccinate a mammal against a targeted disorder.
XX The fusion complexes may be used to suppress an immune response in an
XX animal suffering from an immune disorder e.g. multiple sclerosis, insulin
XX -dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or
XX chronic allergies. The complexes may also be used in the treatment of
XX livestock and pets such as cats and dogs. The MHC fusion complexes can be
XX produced such that they contain a single antigenic peptide including one
XX of known structure, additionally a wide range of peptides can be
XX presented for T cell interaction
XX
XX Sequence 1382 BP; 320 A; 374 C; 404 G; 284 T; 0 U; 0 Other;
XX
Query Match 66.0%; Score 606.2; DB 2; Length 1382;
Best Local Similarity 89.5%; Pred. No. 1.5e-130;
Matches 672; Conservative 0; Mismatches 59; Indels 21; Gaps 1;
Qy 1 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGGTGTG 60
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGGTGTCTGATGGTGTG 65
Qy 61 AGCAGCCCGGAGCTGAGGGCGGAACTCCATCTCTCGCCCTCGCTGAGCAGCCCG 120
Db 66 AGCAGCCCGGAGCTGAGGGCGGAGCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 104
Qy 121 ATCGTGTGTCTCCGCGAGCTGGGACGGAGTGGGGGCTCACTAGTCCCGAGGCTCTGA 180
Db 105 GCTGCTCAGCTGAAATCAACGAAGCTGCTGTCTAGCGAGGGGGGAGCGCGGA 164
Qy 181 GGTGAGGCTCCGAAAGGCATTTCTGTGTCAGTTTCAGGGGAGTGTCTACTACACCAAC 240
Db 165 GGGGAAACTCCGAAAGGCATTTCTGTGTCAGTTTCAGGGGAGTGTCTACTACACCAAC 224
Qy 241 GGGAGCGAGCGCATACGGCTGTGTACACAGATACATCTACACCGGAGGAGTACGTGCCG 300
Db 225 GGGAGCGAGCGCATACGGCTGTGTACACAGATACATCTACACCGGAGGAGTACGTGCCG 284
Qy 301 TACGACAGCGACGTGGGGGAGTACCGCGCGGTACCGAGTGGGGGCGGCGAGCGCGAG 360
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GenCore version 5.1.9
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OM protein - nucleic search, using frame_plus_p2n model

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(without alignments)
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Fgapop 6.0 , Fgapext 7.0
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Searched: 6366136 seqs, 31973710525 residues

Total number of hits satisfying chosen parameters: 12732272

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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ALIGNMENTS

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VERSION AX081281.1 GI:13170131
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Glaichenhaus,N. and Malherbe,L.
TITLE Recombinant proteins and molecular complexes derived therefrom,
JOURNAL analogous to molecules involved in immune responses
PATENT Patent: WO 0109194-A 2 08-FEB-2001;
CENTRE CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)
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2	1255.5	77.5	893	2	AR047947 Sequence
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Best Local Similarity: 85.7%          Mismatches: 24
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DB:                  2                Gaps:       3

US-10-048-116B-6 (1-306) x AR175097 (1-1382)

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QY      21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCAGCTGAA----- 119
QY      41 IleValValSerGlySerTrpAspGlyGlyGlyValGlnPheLysGlyGlyCysTyrThrAsn 80
Db      165 GGGGGAACTCCGAAAGGCAATTCGTGTCAGTTCAGGGCGAGTCTACTACACCAAC 224
QY      81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db      225 GGGACGACGCGCATACGGCTCGTGACCATATCTACAAACCGGGGAGTACGTGCGC 284
QY      101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db      285 TAGCAGCAGCAGTGGCGGAGTACCGCGGTCAGCGGTCAGCGGCGGCGGCGGAG 344
QY      121 TyrTrpAsnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db      345 TACTGGAAACAGCAGCGCGGAGATCTTGGAGCGAACCGGGCGGAGGTGGACACGGCGTGC 404
QY      141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db      405 AGACACAACCTACGAGGGCGGAGACCGACCTCCCTCGCGCGGCTTGAACAGCCCAAT 464
QY      161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db      465 GTCCCATCTCTCTGTCAGGACAGAGGCCCTCAACCAACCAACACTCTGTGTCTGTTCG 524
QY      181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGluGlu 200
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ORIGIN
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Pred. No.:            1145.00          Matches:    227
Score:                87.2%           Conservative: 4
Best Local Similarity: 85.7%          Mismatches: 24
Query Match:         70.7%           Indels:     10
DB:                  2                Gaps:       3

US-10-048-116B-6 (1-306) x AR175097 (1-1382)

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QY      21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCAGCTGCTCAGCTGAA----- 119
QY      41 IleValValSerGlySerTrpAspGlyGlyGlyValGlnPheLysGlyGlyCysTyrThrAsn 80
Db      165 GGGGGAACTCCGAAAGGCAATTCGTGTCAGTTCAGGGCGAGTCTACTACACCAAC 224
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Db      225 GGGACGACGCGCATACGGCTCGTGACCATATCTACAAACCGGGGAGTACGTGCGC 284
QY      101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db      285 TAGCAGCAGCAGTGGCGGAGTACCGCGGTCAGCGGTCAGCGGTCAGCGGCGGCGGAG 344
QY      121 TyrTrpAsnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db      345 TACTGGAAACAGCAGCGCGGAGATCTTGGAGCGAACCGGGCGGAGGTGGACACGGCGTGC 404
QY      141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db      405 AGACACAACCTACGAGGGCGGAGACCGACCTCCCTCGCGCGGCTTGAACAGCCCAAT 464
QY      161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db      465 GTCCCATCTCTCTGTCAGGACAGAGGCCCTCAACCAACCAACACTCTGTGTCTGTTCG 524
QY      181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGluGlu 200
Db      525 GTGACAGATTTCTACCCAGCCCAAGATCAAAGTCGCGTGTTCAGGAATGCCAGGAGGAG 584
QY      201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220

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Db      585 ACATGGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTCCAGGTCCTG 644
Qy      221 ValMetLeuGluMetThrProHisGlnGlyValTyrThrCysHisValGluHisPro 240
Db      645 GTATGCTGGAGATGACCCCTCATCAGGAGAGGTTCTACCTCGCATGTGGAGCATCCC 704
Qy      241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
Db      705 AGCGTGAAGAGCGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy      261 GlyGlyGlyGlySer 265
Db      756 GCGCGTGGTGGTTCC 770

RESULT 8
AX032545
LOCUS   AX032545               1382 bp      DNA          linear      PAT 20-SEP-2000
DEFINITION   Sequence 123 from Patent EP0997477.
ACCESSION   AX032545
VERSION     AX032545.1   GI:10279486
KEYWORDS    .
SOURCE      unidentified
            unidentified
            unclassified sequences.
REFERENCE   1
            Chavallaz, P.A., Edwards, A.C., Grammer, S., Jiao, J.A., Rhode, P.R.,
            Weidanz, J.A. and Wong, H.C.
            Mhc complexes and uses thereof
            Patent: EP 0997477-A 123 03-MAY-2000;
            JOURNAL   SUNOL MOLECULAR CORP (US)
FEATURES    Location/Qualifiers
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             /organism="unidentified"
             /mol_type="unassigned DNA"
             /db_xref="taxon:32644"

ORIGIN
Alignment Scores:
Pred. No.:      1.91e-110      Length:      1382
Score:          1145.00      Matches:      227
Percent Similarity: 87.2%      Conservative: 4
Best Local Similarity: 85.7%      Mismatches: 24
Query Match:    70.7%      Indels:      10
DB:             2           Gaps:         3

US-10-048-116B-6 (1-306) x AX032545 (1-1382)

Qy      1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
Db      6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGTCTGCTGTGGTGTGCTGATGGTGTG 65
Qy      21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      66 AGCAGCCCAAGAGCTTAAAGTATCTCAGGCTGTTCACGCTGCTCAGCTGAA-----119
Qy      41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db      120 ATCAACGAAGCTGGTCTGCTAGCGAGGGGGCGGAGC-----GGCGGA 164
Qy      61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrTyrThrAsn 80
Db      165 GGGGAAACTCCGAAGGCAATTCGTGGTCCAGTTCAGGCTGTTCACGCTGCTCAGCTGAA 224
Qy      81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db      225 GGGAGCAGCGATACCGCTCGTGACCAAGATACATCTCAACCCGGAGGATGACGTGGCG 284
Qy      101 TyrAspSerAspValGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db      285 TACGACGCGACGTGGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCGACGCGAG 344
Qy      121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140

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Db      345 TACTGGAACAGCAGCGAGATCCTGGAGCGAACCGGGCGAGGTGGACACGGCGTGC 404
Qy      141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db      405 AGACAACAATACGAGGGCGGAGACCGACCTCCCTCGCGCGGCTTGAACAGGCCAAT 464
Qy      161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db      465 GTCCGCATCTCCCTGCTCCAGGACAGAGGCCCTCAACACCAACACACTCTGTGTCTGTCG 524
Qy      181 ValThrAspPheTyrProAlaLysIleLysValArgTyrPheArgAsnGlyGlnGluGlu 200
Db      525 GTGACAGATTCTACCCAGCCCAAGATCAAGTGCCTGCTTCAAGAAATGCCAGGAGGAG 584
Qy      201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrPheGlnValLeu 220
Db      585 ACAGTGGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGCTGGACCTTCCAGGTCCTG 644
Qy      221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db      645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTTCTACACCTGCATGTGGAGCATCCC 704
Qy      241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
Db      705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy      261 GlyGlyGlyGlySer 265
Db      756 GCGCGTGGTGGTTCC 770

RESULT 9
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LOCUS   AR033962               1385 bp      DNA          linear      PAT 29-SEP-1999
DEFINITION   Sequence 121 from patent US 5869270.
ACCESSION   AR033962
VERSION     AR033962.1   GI:5949567
KEYWORDS    .
SOURCE      Unknown.
            Unknown.
            Unclassified.
REFERENCE   1 (bases 1 to 1385)
            Rhode, P.R., Jiao, J.-A., Burkhardt, M. and Wong, H.C.
            Single chain MHC complexes and uses thereof
            Patent: US 5869270-A 121 09-FEB-1999;
            JOURNAL   Location/Qualifiers
FEATURES    Location/Qualifiers
             1..1385
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ORIGIN
Alignment Scores:
Pred. No.:      1.91e-110      Length:      1385
Score:          1145.00      Matches:      227
Percent Similarity: 87.2%      Conservative: 4
Best Local Similarity: 85.7%      Mismatches: 24
Query Match:    70.7%      Indels:      10
DB:             2           Gaps:         3

US-10-048-116B-6 (1-306) x AR033962 (1-1385)

Qy      1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
Db      6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGTCTGCTGTGGTGTGCTGATGGTGTG 65
Qy      21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      66 AGCAGCCCAAGAGCTTAAAGTATCTCAGGCTGTTCACGCTGCTCAGCTGAA-----119
Qy      41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db      120 ATCAACGAAGCTGGTCTGCTAGCGAGGGGGCGGAGC-----GGCGGA 164
Qy      61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrTyrThrAsn 80

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Db      165  GGGGAACTCCGAAGGCAATTCGTGTCGATTCAGGGCGAGTGTCTACTACCAAC 224
Qy      81  GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db      225  GGGACGCGAGCGCATACCGCTCGTGACAGATACATCTACACCGGAGGAGTACGTGGC 284
Qy      101  TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db      285  TACACAGCGACGCGGGCGAGTACCGCGCGGTGACCGAGTGGGCGCGCCAGACGCGG 344
Qy      121  TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db      345  TACTGGAAACAGCCGCGAGATCCTGGAGGGAACGCGGGCGGAGGTGGACACGCGTGC 404
Qy      141  ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db      405  AGACACAACCTACGAGGGCGGAGACCGACCTCCCTGCGGGCGCTTGAAACAGCCCAAT 464
Qy      161  ValAlaIleSerLeuSerArgThrGluAlaValThrArgAlaGluValAspThrAlaCys 180
Db      465  GTGCCATCTCCCTGTCCAGGACAGAGGCGCTCAACACCAACACACACTCTGTGTTCG 524
Qy      181  ValThrAspPheTyrProAlaValIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
Db      525  GTACAGATTTCTACCCAGCAAGATCAAAAGTGGCTGTGTTAGGAATGGCCAGGAGAG 584
Qy      201  ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220
Db      585  ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTTG 644
Qy      221  ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db      645  GTATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy      241  SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
Db      705  AGCCTGAAGAGCCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy      261  GlyGlyGlyGlySer 265
Db      756  GCGCGTGGTGGTTC 770

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RESULT 10

AR175095
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source

AR175095.1 GI:17916394

Unknown.

Unknown.

Unclassified.

1 (bases 1 to 1385)

Rhode, P.R., Jiao, J.-A., Burkhardt, M. and Wong, H.C.

MHC molecules and uses thereof

Patent: US 6309645-A 121 30-OCT-2001;

Location/Qualifiers

1..1385

/organism="unknown"

/mol_type="unassigned DNA"

ORIGIN

Alignment Scores:

Pred. No.: 1.91e-110 Length: 1385
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 70.7% Indels: 10
DB: 2 Gaps: 3

US-10-048-116B-6 (1-306) x AR175095 (1-1385)

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Qy      1  MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20
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Qy      21  SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      66  AGCAGCCCAAGGACCTTAAGTATCTCAAGCTGTTCACGCTGTCTACGCTGAA----- 119
Qy      41  IleValValSerGlySerTrpAspGlyGlyGlyGlySerLeuValProArgGlySerGly 60
Db      120  ATCAACGAAGCTGGTGTGTGTAGCGAGGGGGCGGAAGC-----GGCGGA 164
Qy      61  GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrTyrThrAsn 80
Db      165  GGGGGAAACTCCGAAGGCAATTCGTGTGTCAGTTCAGGGCGAGTGTCTACTACCAAC 224
Qy      81  GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db      225  GGGACGCGAGCGCATACCGCTCGTGACAGATACATCTACACCGGAGGAGTACGTGGC 284
Qy      101  TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db      285  TACACAGCGACGCGGGCGAGTACCGCGCGGTGACCGAGTGGGCGCGCCAGACGCGG 344
Qy      121  TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db      345  TACTGGAACAGCCGAGCGGAGATCTCTGGAGCAACCGCGGCGGCTTGAAACAGCCCAAT 464
Qy      141  ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db      405  AGACACAACCTACGAGGGCGGAGACCGACCTCCCTGCGGGCGCTTGAAACAGCCCAAT 464
Qy      161  ValAlaIleSerLeuSerArgThrGluAlaValAsnHisAsnThrLeuValCysSer 180
Db      465  GTGCCATCTCCCTGTCCAGGACAGAGGCGCTCAACACCAACACACTCTGTGTCTGTCG 524
Qy      181  ValThrAspPheTyrProAlaValIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
Db      525  GTACAGATTTCTACCCAGCAAGATCAAAAGTGGCTGTGTTAGGAATGGCCAGGAGAG 584
Qy      201  ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220
Db      585  ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTTG 644
Qy      221  ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db      645  GTATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy      241  SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
Db      705  AGCCTGAAGAGCCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy      261  GlyGlyGlyGlySer 265
Db      756  GCGCGTGGTGGTTC 770

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RESULT 11

CS079299
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source

CS079299.1 GI:63093741

unidentified

unclassified

unclassified sequences.

1

Rhode, P.R., Jiao, J.A., Burkhardt, M. and Wong, H.C.

MHC complexes and uses thereof

Patent: EP 1526141-A 121 27-APR-2005;

Location/Qualifiers

1..1385

/organism="unknown"

/mol_type="unassigned DNA"

unclassified sequences.

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Rhode, P.R., Jiao, J.A., Burkhardt, M. and Wong, H.C.

MHC complexes and uses thereof

Patent: EP 1526141-A 121 27-APR-2005;

Location/Qualifiers

1..1385

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ORIGIN

Alignment Scores:

Pred. No.: 1.91e-110 Length: 1385
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 70.7% Indels: 10
DB: 2 Gaps: 3

US-10-048-116B-6 (1-306) x CS079299 (1-1385)

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Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTG 65
Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGACCTTAAGTATCTCTCAGGCTGTTACGCTGCTCAGCTGAA----- 119
Qy 41 IleValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAAGCTGGTGGTCTAGCGGAGGGGGCGGAAGC-----GGCGGA 164
Qy 61 GlyGlyGlySerGluArgHisPheValGlnPheLysGlyGluCysTyrThrAsn 80
Db 165 GGGGGAACCTCCGAAGGACATTCGTGTGTCAGTTCAGAGGGCGAGTGTACTACACCAAC 224
Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGAGCGAGGATCCGAGGACCTTCTCAGGCTGTTACGCTGCTCAGCTGAA----- 284
Qy 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluGluArgProAspAlaGlu 120
Db 285 TACGACAGCGACGTGGGCGAGTACCGCGGTGACCGAGTGGGCGGCCAGACGCCGAG 344
Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAACTCCGAGGAGATCTGGAGCGAACCGCGGCGGAGGTGGACACCGCGCTGC 404
Qy 141 ArgHisAsnTyrGluGlyProGluThrSerLeuArgArgGluGluGlnProAsn 160
Db 405 AGACACAACTACGAGGCGCGAGACCGACCTCTCTCGGCGGCTTGAACAGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db 465 GTCCCATCTCCCTGTCCAGGACAGAGCCCTCAACACCAACCAACACCTCTGTGTTCG 524
Qy 181 ValThrAspPheTyrProAlaIleLysValArgTrpPheArgAsnGlyGlnGlu 200
Db 525 GTGACAGATTCTACCCAGCAAGATCAAGATGCGCTGTTCAGGAAATGGCCAGAGGAG 584
Qy 201 ThrValGlyValSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220
Db 585 ACAGTGGGGGTCTCATCCACAGACTTATAGGAATGGGGACTGGACCTTCAGGTCTCTG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATCTGGAGATGACCCCTCATCAGGAGAGGTCTACCTGCGCATGTGGAGCATCCC 704
Qy 241 SerLeuValSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
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Qy 261 GlyGlyGlyGlySer 265
Db 756 GGCCTGTGTGTTC 770

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RESULT 12
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LOCUS AX032543 1385 bp DNA linear PAT 20-SEP-2000
DEFINITION Sequence 121 from Patent EP0997477.
ACCESSION AX032543
VERSION AX032543.1 GI:10279484
KEYWORDS .
SOURCE unidentified
ORGANISM unidentified sequences.
REFERENCE 1
AUTHORS Chavallaz,P.A., Edwards,A.C., Grammer,S., Jiao,J.A., Rhode,P.R.,
Weidanz,J.A. and Wong,H.C.
TITLE Mhc complexes and uses thereof
JOURNAL Patent: EP 0997477-A 121 03-MAY-2000;
SUNOL MOLECULAR CORP (US)
FEATURES
source
1..1385
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ORIGIN
Alignment Scores:
Pred. No.: 1.91e-110 Length: 1385
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 70.7% Indels: 10
DB: 2 Gaps: 3

US-10-048-116B-6 (1-306) x AX032543 (1-1385)
Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTG 65
Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGACCTTAAGTATCTCTCAGGCTGTTACGCTGCTCAGCTGAA----- 119
Qy 41 IleValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAAGCTGGTGGTCTAGCGGAGGGGGCGGAAGC-----GGCGGA 164
Qy 61 GlyGlyGlySerGluArgHisPheValGlnPheLysGlyGluCysTyrThrAsn 80
Db 165 GGGGGAACCTCCGAAGGACATTCGTGTGTCAGTTCAGAGGGCGAGTGTACTACACCAAC 224
Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGAGCGAGGATCCGAGGACCTTCTCAGGCTGTTACGCTGCTCAGCTGAA----- 284
Qy 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluGluArgProAspAlaGlu 120
Db 285 TACGACAGCGACGTGGGCGAGTACCGCGGTGACCGAGTGGGCGGCCAGACGCCGAG 344
Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAACTCCGAGGAGATCTGGAGCGAACCGCGGCGGAGGTGGACACCGCGCTGC 404
Qy 141 ArgHisAsnTyrGluGlyProGluThrSerLeuArgArgGluGluGlnProAsn 160
Db 405 AGACACAACTACGAGGCGCGAGACCGACCTCTCTCGGCGGCTTGAACAGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db 465 GTCCCATCTCCCTGTCCAGGACAGAGCCCTCAACACCAACCAACACCTCTGTGTTCG 524
Qy 181 ValThrAspPheTyrProAlaIleLysValArgTrpPheArgAsnGlyGlnGlu 200
Db 525 GTGACAGATTCTACCCAGCAAGATCAAGATGCGCTGTTCAGGAAATGGCCAGAGGAG 584
Qy 201 ThrValGlyValSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220

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Db 585 ACAGTGGGGTCTCATCACACACCTTATTAGGAATGGGACTGGACCTTCCAGGTCTG 644
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Qy 241 SerLeuLysSerProIleThrValGluThrArgAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCTGAGAGAGCCCATCATCTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGCGTGGTGGTTCC 770

RESULT 13
AR033963
LOCUS AR033963 1508 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 122 from patent US 5869270.
ACCESSION AR033963
VERSION AR033963.1 GI:5949568
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1508)
AUTHORS Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
TITLE Single chain MHC complexes and uses thereof
JOURNAL Patent: US 5869270-A 122 09-FEB-1999;
FEATURES
LOCATION/Qualifiers
source 1..1508.
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ORIGIN
Alignment Scores: 2.13e-110 Length: 1508
Pred. No.: 1145.00 Matches: 227
Score: 87.2% Conservative: 4
Percent Similarity: 85.7% Mismatches: 24
Best Local Similarity: 70.7% Indels: 10
Query Match: 2 Gaps: 3
DB:

US-10-048-116B-6 (1-306) x AR033963 (1-1508)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
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Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCACGCTGCTCAGCGTGA----- 119
Qy 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAGCTGCTGCTAGCGAGGGGGGGGAGC-----GGCGGA 164
Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrrThrAsn 80
Db 165 GGGGGAATCTCCGAAAGGCAATTCGTGTGTCCAGTTCAGGGCGAGTGCTACTACACCAAC 224
Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrrIleTyrrAsnArgGluGluThrValArg 100
Db 225 GGGACGACGGCATACGGCTCGTGACCATATCATCTACACCGGGAGGAGTACGTGCGC 284
Qy 101 TyrAspSerAspValGlyGluTyrrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db 285 TACGACAGCGAGCTGGCGAGTACCGCGGGTGACCGAGCTGGGGCGGCAGACCGCCGAG 344
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Db 345 TACTGGAAACGACGCGGAGATCTCTGGAGCGAAACGCGGGCGAGGTGGACCGGGTGC 404
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Db 405 AGACACAACTACGAGGGGCGGAGACAGCACCTCCTCGCGCGGCTTGAACAGCCCAAT 464
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Db 525 GTGACAGATTCTTACCAGCCCAAGATCAAAGTGGCTGGTTTCAGGAATGGCAGAGGAG 584
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Db 585 ACAGTGGGGTCTCATCCACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyValThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
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Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGCGTGGTGGTTCC 770

RESULT 14
AR152030
LOCUS AR152030 1508 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 24 from patent US 6232445.
ACCESSION AR152030
VERSION AR152030.1 GI:15118080
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1508)
AUTHORS Rhode,P.R., Acevedo,J., Burkhardt,M., Jiao,J.-a. and Wong,H.C.
TITLE Soluble MHC complexes and methods of use thereof
JOURNAL Patent: US 6232445-A 24 15-MAY-2001;
FEATURES
LOCATION/Qualifiers
source 1..1508
/organism="unknown"
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ORIGIN
Alignment Scores: 2.13e-110 Length: 1508
Pred. No.: 1145.00 Matches: 227
Score: 87.2% Conservative: 4
Percent Similarity: 85.7% Mismatches: 24
Best Local Similarity: 70.7% Indels: 10
Query Match: 2 Gaps: 3
DB:

US-10-048-116B-6 (1-306) x AR152030 (1-1508)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCAGCCCTCCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTGTG 65
Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTTCACGCTGCTCAGCGTGA----- 119
Qy 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAGCTGCTGCTAGCGAGGGGGGGGAGC-----GGCGGA 164
Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrrThrAsn 80
Db 165 GGGGGAATCTCCGAAAGGCAATTCGTGTGTCCAGTTCAGGGCGAGTGCTACTACACCAAC 224
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Qy 81 GlyThrGlnArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGAGCGCAGCGATACGGCTCTGACCAAGATACATCTACACCGGAGGAGTACGTGGC 284
Qy 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db 285 TACGACAGCGACGTGGCGAGTACCGCGGTGACCGAGCTGGGCGGCCACAGCCCGAG 344
Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTCGAACAGCAGCCGAGATCCTGGAGCGAACCGCGCGCGAGGTGACACGCGCTGC 404
Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgGluGlnProAsn 160
Db 405 AGACACACTAGAGGGGCGGAGACCGACCTCTCCGCGCGCTTGAACAGGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrIleuValCysSer 180
Db 465 GTCCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCAACACACTCTGGTCTGTTGC 524
Qy 181 ValThrAspPheTyrProAlaIleValIleValArgTyrPheArgAsnGlyGlnGlu 200
Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGCCTGTTTCAGGAATGGCCAGGAG 584
Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrPheGlnValLeu 220
Db 585 ACAGTGGGGTCTCATCCACAGCTTATTAGGAATGGGACTGGACCTTCCAGTCTCTG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCCTGAAGAGAGCCCATCACTGTGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGCGTGGTGTTC 770

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RESULT 15
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 LOCUS ARI75096 1508 bp DNA linear PAT 17-DEC-2001
 DEFINITION Sequence 122 from patent US 6309645.
 ACCESSION ARI75096
 VERSION ARI75096.1 GI:17916395
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 Unclassified.

REFERENCE 1 (bases 1 to 1508)
 AUTHORS Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
 TITLE MHC molecules and uses thereof
 JOURNAL Patent: US 6309645-A 122 30-OCT-2001;
 FEATURES
 Location/Qualifiers
 1..1508
 /organism="unknown"
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ORIGIN
 Alignment Scores:
 Pred. No.: 2,13e-110 Length: 1508
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116B-6 (1-306) x ARI75096 (1-1508)

```

Qy 1 MetAlaLeuGlnIleProSerIleuLeuSerAlaAlaValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCCCAGCGCTCCTCTCAGCTGTGTTGGTCTGATGGTGTCTG 65

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Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerSerLeuGluHisPro 40
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Qy 41 IleValValSerGlySerTrpAspGlyGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAGAGCTGGTGGTCTAGCGAGGGGGGGGAGC-----GGCGGA 164
Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyCysTyrTyrThrAsn 80
Db 165 GGGGAAACTCCGAAAGGCATTTCTGCTGCAGTTCAAGGGCGAGTGTCTACTACCAAC 224
Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGAGCAGCGCATACGGCTCTGACCAAGATACATCTCAACCGGAGGAGTACGTGGC 284
Qy 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db 285 TACGACAGCGACGTGGCGAGTACCGCGCGGTGACCGAGCTGGGCGGCCACAGCCCGAG 344
Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAACAGCCAGCCGAGATCCTGGAGCGAACCGCGCGAGGTGACACCGCGTGC 404
Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db 405 AGACACACTACGAGGGGCGGAGACCGACCTCTCCGCGCGCTTGAACAGGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrIleuValCysSer 180
Db 465 GTCCGCATCTCCCTGTCCAGGACAGAGGCCCTCAACACCAACACACTCTGGTCTGTTGC 524
Qy 181 ValThrAspPheTyrProAlaIleValIleValArgTyrPheArgAsnGlyGlnGlu 200
Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGCCTGTTTCAGGAATGGCCAGGAG 584
Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrPheGlnValLeu 220
Db 585 ACAGTGGGGTCTCATCCACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTCTG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCCTGAAGAGAGCCCATCACTGTGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGCGTGGTGTTC 770

```

Search completed: June 30, 2006, 05:17:54
 Job time : 6500.66 secs

GenCore version 5.1.9
Copyright (c) 1993 - 2006 Bioceleration Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: June 30, 2006, 01:23:04 ; Search time 532.723 Seconds
(without alignments)
6007.369 Million cell updates/sec

Title: US-10-048-116B-6
Perfect score: 1620
Sequence: 1 MALQPSLLLSAAVVVLMVL.....LKWKLQALKKKLQAQHHHHH 306

Scoring table: BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 5244920 seqs, 3486124231 residues

Total number of hits satisfying chosen parameters: 10489840

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlp
-Q=/abes/ABSSWEB spool/US10048116/runat_29062006_093309_10102/app_query.fasta_1
-DB=N Geneseq -QFMT=fastap -SUFFIX=p2n.Tng -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
-UNITS=bites -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
-DOALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=ptc -NORM=ext -HEAPSIZ=500 -MINLEN=0 -MAXLEN=200000000 -HOST=abes05p
-USER=US10048116 @CGN 1.1 1423 @runat_29062006_093309_10102 -NCPU=6 -ICPU=3
-NO MAP -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG -DEV TIMEOUT=120
-WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

N_Geneseq 8:
1: Geneseqn1980s:
2: Geneseqn1990s:
3: Geneseqn2000s:
4: Geneseqn2001as:
5: Geneseqn2001bs:
6: Geneseqn2002as:
7: Geneseqn2002bs:
8: Geneseqn2003as:
9: Geneseqn2003bs:
10: Geneseqn2003cs:
11: Geneseqn2003ds:
12: Geneseqn2004as:
13: Geneseqn2004bs:
14: Geneseqn2005s:
15: Geneseqn2006s:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1620	100.0	921	5 AAF55099	Aaf55099 DNA encod
2	1255.5	77.5	893	2 AAT04262	Aat04262 Hybrid IA
3	1235	76.2	945	12 ADQ31225	Adq31225 I-Ab(beta

ALIGNMENTS

RESULT 1

ID AAF55099 standard; DNA; 921 BP.

XX AAF55099;

XX 15-MAY-2001 (first entry)

XX DNA encoding a fusion protein comprising a beta chain of MHC.

XX Recombinant protein; alpha chain; beta chain; MHC; immunoglobulin;
KW major histocompatibility complex; Fc region; antigen; T lymphocyte;
KW immunostimulant; vaccine; infection; tumour; ss.

OS Synthetic.

XX Key Location/Qualifiers

FT CDS 1..921

FT /*tag= a

XX WO200109194-A1.

XX 08-FEB-2001.

XX 28-JUL-2000; 2000WO-FR002193.

XX 29-JUL-1999; 99FR-00009862.

Adq31228 I-Ab(beta
Aav12068 Murine IA
Aat04269 Hybrid IA
Aat17588 Vector SC
Aat86989 SCEL sing
Aca60744 Mouse MHC
Aat17586 Vector SS
Aat86987 SSCI sing
Aca60742 Mouse MHC
Aat17587 Vector SC
Aat86988 SCTL sing
Aax89069 Single ch
Aca60743 Mouse MHC
Adj75986 Marker ge
Adx26090 Novel cel
Abi99040 Murine pc
Abi99038 Murine pc
Abi99039 Murine pc
Aaq03170 Sequence
Aat06286 I-Ab-beta
Aaq56920 Mouse I-A
Abn84048 Single ch
Aaq35055 IAB beta
Abi99031 MBP 1-14
Abi99028 IAS MBP 1
Abi99032 MBP 1-14
Abi99027 IAS MBP 1
Abi99030 IAS MBP 9
Abi99021 I-Ae MBP.
Abi99029 IAS MBP 9
Abi99033 MBP 90-10
Aat60705 cDNA enco
Aat60700 cDNA enco
Aec64482 DRB1-biot
Aad63150 Human maj
Aad62751 Human maj
Adp88246 Lung canc
Adr24869 Breast ca
Ado40822 DNA encod
Abk63510 Rat seque
Adb57995 Toxicity-
Abt41775 Toxicity

ADQ31228
AAV12068
AAT04269
AAT17588
AAT86989
ACA60744
AAT17586
AAT86987
ACA60742
AAT17587
AAT86988
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AAT06286
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ABI99027
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1096.5 67.7 798
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1049.5 64.8 1698
1044.5 64.5 1662
979.5 60.5 702
979.5 60.5 702
972 60.0 1243
963.5 59.5 702
957 59.1 1686
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957 59.1 2059
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952 58.8 1707
949 58.6 1680
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854.5 52.7 1323
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839.5 51.8 1192
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839.5 51.8 1192
833.5 51.5 941
829 51.2 562
829 51.2 562
829 51.2 562

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XX Kappler JW, Marrack P;
 XX WPI; 1995-320543/41.
 DR P-PSDB; AAR82538.
 XX
 PT Peptide-MHC complex comprising antigenic peptide, linker and MHC segment
 PT - useful as reagents for the treatment of diseases including auto-immune
 PT diseases, immuno-stimulatory diseases or graft-host rejection.
 XX
 PS Example 2; Page 65; 94pp; English.

XX This sequence represents a hybrid IA beta chain gene. This sequence
 CC contains a fragment of the IE alpha chain (residues 56-73), as well as a
 CC linker and cleavage site. This sequence was transfected into a B cell
 CC line (M12.C3) using plasmid pM12-IAb-Ea. It was found that the encoded
 CC sequence was expressed in these cells. Complexes such as this may be used
 CC to regulate an immune response. The complexes are capable of being
 CC recognised by a TCR alone or in combination with additional MHC proteins.
 CC These complexes are useful for therapeutic purposes and experimental
 CC purposes. They can also be used as reagents for the treatment of diseases
 CC including autoimmune diseases, immunodeficiency diseases,
 CC immunoproliferation diseases, and graft-host rejection

XX Sequence 1013 BP; 220 A; 272 C; 327 G; 192 T; 0 U; 2 Other;

Alignment Scores:

Pred. No.: 2,31e-99 Length: 1013
 Score: 1151.00 Matches: 230
 Percent Similarity: 86.8% Conservative: 6
 Best Local Similarity: 84.6% Mismatches: 22
 Query Match: 71.0% Indels: 14
 DB: 2 Gaps: 4

US-10-048-116B-6 (1-306) x AAT04269 (1-1013)

QY 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db 63 ATGGCTCTGAGATCCCAAGCCCTCCCTCTCGGCTGCTGGTGGTGCATGGTGGCTG 122
 QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 123 AGCAGCCCGGACTGAGGGCGAGACTCC-----GAAGCTAGCTTTGAGGCTCAG 173
 QY 41 -----tLeValValSerGlySerTrpAspGlyGlyGlyGlySerLeuVal 55
 Db 174 GGTGCACCTGGCCCAACATTCGTGCGACAGGCTGGAGTGGTGGTCCGCTGGA----- 227
 QY 56 ProArgGlySerGlyGlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlu 75
 Db 228 ---GGGGGAAGTGGAGGTGGAGGGTCTGAAAGGCATTCGTGPACCACTTCATGGCGGAG 284
 QY 76 CysTyrTyrThrAsnGlyThrGlnArgIleAtqLeuValThrArgTyrIleTyrAsnArg 95
 Db 285 TGCTACTTACCACCGGGACGCGCCGACGACGATATGTGACCGAGATACATCTCAACACCGG 344
 QY 96 GluGluTyrValArgTyrAspSerAspValGlyGlyTyrArgAlaValThrGluLeuGly 115
 Db 345 GAGGAGTACGTGGCTACGACAGCGAGCGTGGGGGAGCAGCGCGGTGACCGAGCTGGGG 404
 QY 116 ArgProAspAlaGluTyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGlu 135
 Db 405 CGGCCAGAGCCCGAGTACTTGGAAACAGCCAGCCGAGATCCTGGAGCGAAGCGCGGCGGAG 464
 QY 136 ValAspThrAlaCysArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArg 155
 Db 465 GTGGACACCGGTGTCAGACACAACTACAGGGGCCCGAGACCCACACCTCCCTGCCGGCGG 524
 QY 156 LeuGluGlnProAsnValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsn 175
 Db 525 CTTGAACAGCCCAATGTGCTCATCTCCCTGTCAGGACAGAGGCCCTCAACACCCACACAC 584
 QY 176 ThrLeuValCysSerValThrAspPheTyrProAlaLysIleLysValArgTrpPheArg 195

Db 585 ACTCTGCTGCTCAGTGACAGATTCTTACCACGCAAGATCAAGTCCGCTGCTTCGG 644
 QY 196 AsnGlyGlnGluGluThrValGlyValSerSerThrGlnLeuIleAtqAsnGlyAspTrp 215
 Db 645 AATGCCAGAGGAGGACGGTGGGCTCTCATCCACAGCTTATTAGGAATGGGACTGG 704
 QY 216 ThrPheGlnValLeuValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCys 235
 Db 705 ACCTTCAGGTCTCTGCTGAGATGACCCCTCGCGGGGAGAGGTCTAYACCTGT 764
 QY 236 HisValGluHisProSerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGlu 255
 Db 765 CAGCTGGAGATCCACGCTCAAGAGCCCATCATCTGGAGTGGAGGCGACAGTCTGAG 824
 QY 256 SerAlaArgSerLys-----GlyGlyGlyGly 264
 Db 825 TCTGCTGGAGCAAGATGTTGAGCGGCATCGGGGCG 860
 RESULT 7
 AAT17588
 ID AAT17588 standard; DNA; 1382 BP.
 XX
 AC AAT17588;
 XX
 DT 26-SEP-1996 (first entry)
 XX
 DE Vector SCE1-derived single chain gene encoding MHC fusion complex.
 XX
 KW MHC; major histocompatibility complex; PCR; polymerase chain reaction;
 KW T cell activity modulator; antagonist; immune disorder; allergy;
 KW multiple sclerosis; insulin-dependent diabetes mellitus;
 KW rheumatoid arthritis; myasthenia gravis; ds.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 CDS 6..1382
 FT /*tag= a
 FT sig_peptide 6..86
 FT /*tag= b
 FT /label= I-Ad beta chain leader
 FT /note= "murine MHC class II I-Ad gene beta chain leader
 FT sequence"
 FT misc_feature 87..137
 FT /*tag= c
 FT /label= OVA 323-339
 FT /note= "chicken ovalbumin residues 323-339"
 FT misc_feature 138..167
 FT /*tag= d
 FT /note= "10 residue linker peptide"
 FT misc_feature 168..452
 FT /*tag= e
 FT /label= I-Ad beta1
 FT /note= "murine MHC class II I-Ad gene beta-1 domain"
 FT misc_feature 453..734
 FT /*tag= f
 FT /label= I-Ad beta2
 FT /note= "murine MHC class II I-Ad gene beta-2 domain"
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 FT /*tag= h
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 FT /*tag= i
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 FT misc_feature 1353..1379
 FT /*tag= j
 FT /note= "EE tag"

XX PN WO9604314-A1.
 XX PD 15-FEB-1996.
 XX PF 31-JUL-1995; 95WO-US009816.
 XX PR 29-JUL-1994; 94US-00283302.
 XX PR 01-FEB-1995; 95US-00382454.
 XX PA (DADE-) DADE INT INC.
 XX PI Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
 XX PI Chavallaz P, Jiao J;
 XX DR WPI: 1996-129343/13.
 XX DR P-PSDB; AAR98907.
 XX
 PT Major histocompatibility complex fusion complex for modulating T cell
 PT activity - used in the treatment of immune disorders, e.g. multiple
 PT sclerosis, IDDM and rheumatoid arthritis.
 XX PS Example 17; Fig 29; 210pp; English.
 XX
 CC AAT17588 encodes a murine MHC fusion complex capable of modulating T cell
 CC activity encoded by the vector SCE1. The MHC fusion complex comprises at
 CC least one MHC molecule containing a peptide-binding groove and a
 CC presenting peptide covalently linked to the MHC molecule and opt. a
 CC transmembrane domain. DNA encoding a MHC fusion complex may be cloned
 CC into a host cell to express the complex. The transformed cells may then
 CC be used to identify peptides that modulate, pref. antagonise, T cell
 CC activity. DNA encoding a MHC fusion complex or a single chain fusion
 CC molecule may be used to vaccinate a mammal against a targeted disorder.
 CC The fusion complexes may be used to suppress an immune response in an
 CC animal suffering from an immune disorder e.g. multiple sclerosis, insulin
 CC dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or
 CC chronic allergies. The complexes may also be used in the treatment of
 CC livestock and pets such as cats and dogs. The MHC fusion complexes can be
 CC produced such that they contain a single antigenic peptide including one
 CC of known structure, additionally a wide range of peptides can be
 CC presented for T cell interaction
 XX
 SQ Sequence 1382 BP; 320 A; 374 C; 404 G; 284 T; 0 U; 0 Other;
 Alignment Scores:
 Align. No.: 1.3e-98 Length: 1382
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 2 Gaps: 3
 US-10-048-116B-6 (1-306) x AAT17588 (1-1382)
 Qy 1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValMetValLeu 20
 Db 6 ATGCTGTCAGATCCCGAGCTCTCTCTCAGCTGCTGCTGCTGCTGCTGCTGCTG 65
 Qy 21 SerSerProGlyThrGluGlyGlyAenSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTACGCTGCTCAGCTGAA----- 119
 Qy 41 IleValValSerGlySerTipAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db 120 ATCAACGAGCTGGTCTGCTAGCGAGGGGGCGGAAC-----GGCGGA 164
 Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLeuGlyGlyCysTyrTyrThrAsn 80
 Db 165 GGGGAAACTCCGAAGGCAATTCGTGCTCAGTTCAGGGCGAGTGCTACTACACCAAC 224
 Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 Db 225 GGGACGCGAGCATACCGCTCTCGTACCAGATACATCTTAAACCGGGAGGAGTACGTGGC 284

Qy 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db 285 TACGACGCGAGCTGGCGAGTACCGCGGGTACCGAGCTCGGGCGGCACACGCCGAG 344
 Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 Db 345 TACTGGACACGCCAGCGGAGATCTCTGGAGCGAACCGGGCCGAGGTGGACACGGCGTGC 404
 Qy 141 ArgHisAsnTyrGluGlyProGluThrSerLeuArgArgLeuGluGlnProAsn 160
 Db 405 AGACACAACCTACGAGGGCGGAGACACGACCTCTCTCGCGCGCTTGAACAGCCCAAT 464
 Qy 161 VallalIleSerLeuSerArgThrGluAlaLeuAenHisAenThrLeuValCysSer 180
 Db 465 GTCCGCATCTCTCTGTCAGGACAGAGGCCCTCAACACCACCAACACTCTGGTCTGTCG 524
 Qy 181 ValThrAspPheTyrProAlaLysIleLysValArgTppPheArgAsnGlyGlnGluGlu 200
 Db 525 GTGACAGATTTCTACCCAGCCCAAGATCAAGTGGCTGCTGTTCCAGGAATGGCCAGGAGGAG 584
 Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrThrPheGlnValLeu 220
 Db 585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGGACTGGACCTTCCAGGTCTCTG 644
 Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACCTGCATGTGGAGGATCC 704
 Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 Db 705 AGCCTGAAGAGCCCATCATCTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
 Qy 261 GlyGlyGlyGlySer 265
 Db 756 GCGCGTGGTGGTTC 770
 RESULT 8
 ID AAT86989 standard; DNA; 1382 BP.
 AC AAT86989;
 XX 27-MAR-1998 (first entry)
 DE SCE1 single chain gene.
 KW Construction; major histocompatibility complex; MHC; fusion complex;
 XX SCE1 single chain gene; ss.
 OS Synthetic.
 FH Key Location/Qualifiers
 FT CDS 6..1382
 FT /*tag= a
 XX
 XX WO9728191-A1.
 PD 07-AUG-1997.
 XX 30-JAN-1997; 97WO-US001617.
 XX 31-JAN-1996; 96US-00596387.
 XX (DADE-) DADE INT INC.
 XX Rhode PR, Jiao J, Burkhardt M, Wong HC;
 XX WPI: 1997-402555/37.
 XX P-PSDB; AAN29214.
 XX Single chain major histocompatibility complex comprising linked alpha and
 XX beta chains - useful for suppressing an immune response to an auto-immune

PT disease, e.g. multiple sclerosis, rheumatoid arthritis, diabetes
 XX mellitus, etc.
 XX
 XX Example 17; Page 140-141; 217pp; English.
 XX
 CC The present sequence was used in the construction of major
 CC histocompatibility complex (MHC) fusion complexes
 XX
 SQ Sequence 1382 BP; 320 A; 373 C; 405 G; 284 T; 0 U; 0 Other;

 Alignment Scores:
 Pred. No.: 1,3e-98 Length: 1382
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 2 Gaps: 3

 US-10-048-116B-6 (1-306) x AAT86989 (1-1382)

 QY 1 MetAlaLeuGlnIleProSerLeuLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 |||||
 Db 6 ATGGCTCTCGAGATCCCAAGCCTCTCTCTCTCAGCTGCTGTGTGTGTGTGTGTGTGTGTG 65
 |||||
 QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 |||||
 Db 66 AGCAGCCCAAGGACCTTAAGTATCTCTCAGGCTGTTCACGCTGCTCACGCTGAA----- 119
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 QY 41 IleValValSerGlySerTrpAspGlyGlyGlyGlySerLeuValProArgGlySerGly 60
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 Db 120 ATCAACGAAGCTGTGTGTGTCTAGCGAGGGGGCGGAAGC-----GGCGGA 164
 |||||
 QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheIleGlyGlyGlyGluCysTyrThrAsn 80
 |||||
 Db 165 GGGGGAAATCCGAAGAAGCATTTCTGTTCAGTTCAAGGGCGAGTGTCTACTACCCAAC 224
 |||||
 QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 |||||
 Db 225 GGGAGCGACGCATACGGCTCGTACACAGATACATCTACAACCGGGAGGAGTACGTGCGC 284
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 QY 101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
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 Db 285 TAGCAGACGCGAGTGGCGAGTACCGCGCGGTGACCGAGCTGGGGCGGCCAGACGCCGAG 344
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 |||||
 Db 345 TACTGGAAACAGCCCGGAGATCTTGAGAGCAACCGCGGCTGAGGTGGACACGGCGTGC 404
 |||||
 QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
 |||||
 Db 405 AGACACAACCTACGAGGGGCGGAGACCAAGCACCTCCCTCGCGGCGCTTGAACAGCCCAAT 464
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 QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
 |||||
 Db 465 GTCGCCATCTCCCTGTCCAGGACAGAGCCCTCAACACCAACACACTCTGGTGTGTTCG 524
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 QY 181 ValThrAspPheTyrProAlaIleIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
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 Db 525 GTGACAGATTCTTACCCAGCCAAAGATCAAGTGGCTGTGTTCAGAAATGGCCAGGAGGAG 584
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 QY 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
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 Db 585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGAGTGGACCTTCCAGGTCTCTG 644
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 QY 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
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 Db 645 GTCATGCTGGAGATACCCCTCATCAGGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
 |||||
 QY 241 SerLeuLysSerProIleThrValGlnTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 |||||
 Db 705 AGCCTGAAGAGCCCCATCACTGTGAGTGG-----ACTAGTGTGTGGGTGGCAGC 755
 |||||
 QY 261 GlyGlyGlyGlySer 265

agonist and is covalently linked to the MHC protein, or DNA sequence coding for the fusion complex which is a single chain fusion molecule. The methods are useful for identifying a peptide that can modulate the activity of T cells, inducing an immune response in a mammal (including vaccinating a mammal against a targeted disorder) and for suppressing an immune response in a mammal. The disorders include an autoimmune disorder such as multiple sclerosis, insulin-dependent diabetes mellitus, rheumatoid arthritis, myasthenia gravis or chronic allergies. The present sequence encodes a mouse MHC class II I-Ad fusion complex of the invention

Db	705	AGCCTGAAGAGCCCCATCACTGTGAGTGG-----ACTAGTGGTGGCGGTGCAGC	755
Qy	261	GlyGlyGlyGlySer 265 	
Db	756	GGCGGTGGTGGTTCC 770	
RESULT 10			
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ID	AA17586	standard; DNA; 1385 BP.	
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AC	AA17586;		
XX			
DT	26-SEP-1996	(first entry)	
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DE		Vector SSC1-derived single chain gene encoding MHC fusion complex.	
XX			
KW		MHC; major histocompatibility complex; PCR; polymerase chain reaction;	
KW		T cell activity modulator; antagonist; immune disorder; allergy;	
KW		multiple sclerosis; insulin-dependent diabetes mellitus;	
KW		rheumatoid arthritis; myasthenia gravis; ds.	
XX			
OS		Synthetic.	
XX			
Key		Location/Qualifiers	
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FT	sig_peptide	6..86	
FT		/*tag= b	
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FT	misc_feature	138..167	
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FT		/note= "10 residue linker peptide"	
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FT		/*tag= e	
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FT		/note= "murine MHC class II I-Ad gene beta-1 domain"	
FT	misc_feature	453..734	
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FT		/note= "murine MHC class II I-Ad gene beta-2 domain"	
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FT		/*tag= h	
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PN	W09604314-A1.		
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PD	15-FEB-1996.		
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PF	31-JUL-1995;	95WO-US009816.	
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PR	29-JUL-1994;	94US-00283302.	
PR	01-FEB-1995;	95US-00382454.	
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PA	(DADE-) DADE INT INC.		
PI	Wong HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;		

Alignment Scores:

Pred. No.: 1,3e-98 Length: 1385
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 8 Gaps: 3

US-10-048-116B-6 (1-306) x ACA60742 (1-1385)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValLeuMetValLeu 20
 Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGCTGGTGGTGGTGGTGGTGGT 65
 Qy 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGCAGCCCAAGGACTTAAAGTATCTCTCAGGCTGTTACGCTGCTCAGCTGAA----- 119
 Qy 41 IleValValSerGlySerTrpAspGlyGlyGlyGlyGlyGlyGlyGlyGlyGlyGly 60
 Db 120 ATCAACGAAGCTGTGTCTGTAGCGGAGGGGGCGGAAGC-----GGCGGA 164
 Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrTyrThrAsn 80
 Db 165 GGGGAACCTCCGNAAGCAATTCGTGTCTCAGTTCAAGGGCGAGTGTCTACTACCAAC 224
 Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 Db 225 GGGACGCGCGCATACGGCTCGTGACGAGATACATCTACACCGGGAGGAGTACGTGCG 284
 Qy 101 TyrAspSerAspValGlyGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db 285 TAGACAGCGACGTGGCGAGTACCGCGCGGTGACCGAGTGGGGCGCGCGAGCGCGAG 344
 Qy 121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValAspThrAlaCys 140
 Db 345 TACTGGNACAGCAGCGCGAGATCTCGAGCGAACCGCGCGCGAGTGGACACGGCGTGC 404
 Qy 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
 Db 405 AGACACAATACGAGGGCGGAGACCGAGCACCTCCCTCGCGCGCTTGAACAGCCCAAT 464
 Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer 180
 Db 465 GTCCCATCTCTCCGTGTCCAGGACAGAGCGCCCTCAACACCAACCAACACTCTGTGTTC 524
 Qy 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
 Db 525 GTGACAGATTCTACCCAGCAAGATCAAGTGGCTGGTTCAGGAATGCCAGGAGGAG 584
 Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
 Db 585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTTG 644
 Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
 Db 645 GTCATGCTGGAGATGACCCCTCATCGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
 Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 Db 705 AGCCTGAGAGCCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
 Qy 261 GlyGlyGlyGlySer 265
 Db 756 GCGCGTGGTGTTC 770

RESULT 13

AAT17587
 ID AAT17587 standard; DNA; 1508 BP.
 XX
 AC AAT17587;
 XX
 DT 26-SEP-1996 (first entry)

XX DE Vector SCTL-derived single chain gene encoding MHC fusion complex.
 XX KW MHC; major histocompatibility complex; PCR; polymerase chain reaction;
 KW T cell activity modulator; antagonist; immune disorder; allergy;
 KW multiple sclerosis; insulin-dependent diabetes mellitus;
 KW rheumatoid arthritis; myasthenia gravis; ds.
 XX OS Synthetic.
 XX FH Key Location/Qualifiers
 FT CDS 6..1508
 FT /*tag= a
 FT sig_peptide 6..86
 FT /*tag= b
 FT /label= I-Ad beta chain_leader
 FT /note= "murine MHC class II I-Ad gene beta chain leader
 FT sequence"
 FT misc_feature 87..137
 FT /*tag= c
 FT /label= OVA 323-339
 FT /note= "chiGen ovalbumin residues 323-339"
 FT misc_feature 138..167
 FT /*tag= d
 FT /note= "10 residue linker peptide"
 FT misc_feature 168..452
 FT /*tag= e
 FT /label= I-Ad beta1
 FT /note= "murine MHC class II I-Ad gene beta-1 domain"
 FT misc_feature 453..734
 FT /*tag= f
 FT /label= I-Ad beta2
 FT /note= "murine MHC class II I-Ad gene beta-2 domain"
 FT misc_feature 735..806
 FT /*tag= g
 FT /note= "24 residue peptide linker"
 FT misc_feature 807..1067
 FT /*tag= h
 FT /label= I-Ad alpha1
 FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
 FT misc_feature 1068..1352
 FT /*tag= i
 FT /label= I-Ad alpha2
 FT /note= "murine MHC class II I-Ad gene alpha-2 domain"
 FT misc_feature 1353..1505
 FT /*tag= j
 FT /label= I-Ad alpha-TM
 FT /note= "murine MHC class II I-Ad gene alpha-transmembrane
 FT domain"
 XX WO9604314-A1.
 XX PN 15-FEB-1996.
 XX PD 31-JUL-1995; 95WO-US0009816.
 XX PF 29-JUL-1995; 94US-00283302.
 XX PR 01-FEB-1995; 95US-00382454.
 XX PA (DADE-) DADE INT INC.
 XX WONG HC, Rhode PR, Weidanz JA, Grammer S, Edwards AC;
 PI Chavallaz P, Jiao J;
 XX WPI; 1996-129343/13.
 DR P-PSDB; AAR98906.
 XX Major histocompatibility complex fusion complex for modulating T cell
 PT activity - used in the treatment of immune disorders, e.g. multiple
 PT sclerosis, IDDM and rheumatoid arthritis.
 XX Example 17; Fig 28; 210pp; English.
 PS
 XX

QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGCAGCCCCAGACCTTAAGTATCTCTCAGGCTGTTCACGCTGCTACGCTGAA-----119
 QY 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db 120 ATCAACGAAGCTGGTCTGTAGCGAGGGGGGAGC-----GGCGGA 164
 QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrThrAsn 80
 Db 165 GGGGGAACCTCCGAAGGCAITTCGTGTCCAGTTCGAAGGGCGAGTCTACTACACCAAC 224
 QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
 Db 225 GGGAGCGAGCGCATACGCTCGTACACAGATACATCTACAACCGGGAGGAGTACGTGGC 284
 QY 101 TyrAspSerAspValGlyGlyArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db 285 TACGACAGCGAGTGGGCGAGTACCGCGGTGACCGAGCTGGGGGCGGACAGCGCGAG 344
 QY 121 TyrTrpAsnSerGlnProGluLeuGluArgThrArgAlaGluValAspThrAlaCys 140
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 QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
 Db 405 AGACACAACTACGAGGGGCGGAGACCAAGCACCTCCCTGCGGCGCTTGAACAGCCCAAT 464
 QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisAsnThrLeuValCysSer 180
 Db 465 GTCCGCATCTCCTGTCCAGACAGAGGCGCTCAACCAACCAACACTCTGTGTCTGTTCG 524
 QY 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
 Db 525 GTGACAGATTCTACCCAGCCAGCATCAAGTGGCTGGTTCCAGGAATGGCCAGGAGAG 584
 QY 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
 Db 585 ACATGTGGGGTCTCATCCACAGCTTATTAGGAATGGGACTGGACCTTCAGGTCTGT 644
 QY 221 ValMetLeuGluMetThrProHisGlnGlyGlyValTyrThrCysHisValGluHisPro 240
 Db 645 GTCATGTGGAGATGACCCCTCATCAGGAGAGGTCTACACTGCCATGTGGACATCCC 704
 QY 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 Db 705 AGCCTGAAGAGCCCATCACTGTGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
 QY 261 GlyGlyGlySer 265
 Db 756 GCGGTGGTGGTTCC 770

RESULT 15
 AAX89069
 ID AAX89069 standard; DNA; 1508 BP.
 XX
 AC AAX89069;
 XX
 DT 14-SEP-1999 (first entry)
 XX
 DE Single chain IAD/OVA 323-229 MHC fusion protein encoding DNA.
 XX
 KW Major histocompatibility complex; MHC; single chain MHC; sc-MHC; Ig;
 KW peptide binding groove; immunoglobulin; T cell receptor; immune response;
 KW immune-related disorder; antigenic peptide; fusion protein; ss.
 XX
 OS Synthetic.
 XX
 PN WO9921572-A1.
 XX
 PD 06-MAY-1999.
 XX
 PF 13-OCT-1998; 98WO-US021520.

XX

PR

XX

PA

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PI

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DR

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DR

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PT

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PS

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CC

CC

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CC

CC

CC

CC

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CC

CC

CC

CC

CC

SQ

XX

Alignment Scores:

Pred. No.: 1.46e-98 Length: 1508
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservative: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 70.7% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116b-6 (1-306) x AAX89069 (1-1508)

QY

Db

QY

Db

QY

Db

QY

Db

QY

Db

QY

Db

QY

Db

QY

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Db

QY

Db

QY

Db

QY

Db

QY

Db

QY

Db

QY

Db

QY

Db

29-OCT-1997; 97US-00960190.

(SUNO-) SUNOL MOLECULAR CORP.

Rhode PR, Acevedo J, Burkhardt M, Jiao J, Wong HC;

WPI; 1999-418411/35.

P-PSDB; AAY27111.

Single chain major histocompatibility complex class I complexes.

Example 1; Fig 1; 148pp; English.

The invention relates to new single chain major histocompatibility

complex (sc-MHC) class II complexes that comprise a peptide binding

groove, and a modified class II beta 2 chain or covalently linked

immunoglobulin (Ig) light chain constant (C1) region. The MHC complexes

are useful for detection and analysis of peptide ligands, pathogenic T-

cells, for functional, cellular and molecular assays. They can be used to

identify and isolate T cell receptor and/or MHC agonists and antagonists.

They can be used in vivo to compete with pathogenic antigen presenting

cells involved in immune-related disorders. They can also be used to

raise antibodies and to screen immune cells. It is also use in a method

of suppressing an immune response in mammals. The sc-MHC complexes

comprising modified class II beta 2 chains and/or Ig-C1 regions are

soluble and provide enhanced yield. These MHC complexes also can contain

single antigenic peptides readily isolated from expressing cells in

significant quantities. The polyepitopic MHC complexes also provide a

means to detect cells expressing multiple target structures with a single

complex. The present sequence represents a DNA encoding a single chain

IAD/OVA 323-229 MHC fusion protein

SQ Sequence 1508 BP; 337 A; 413 C; 441 G; 317 T; 0 U; 0 Other;

XX

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XX

XX

XX

XX

XX

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XX

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QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db |||||
QY 405 AGACACAACTACAGGGGCGGAGACAGACACTCTCCCTGCGCGGCTTGACAGCCCAAT 464
Db |||||
QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db |||||
QY 465 GTGCCCATCTCCCTGTCCAGACAGAGGCCCTCAACCCACCAACACTCTGGTCTGTTCG 524
Db |||||
QY 181 ValThrAspPheTyrProAlaLysIleLysValArgTrpPheArgAsnGlyGlnGluGlu 200
Db |||||
QY 525 GTGACAGATTTCACCCAGCCAAAGATCAAAGTGCCTGGTTCAGGAATGGCCAGGAGGAG 584
Db |||||
QY 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpThrPheGlnValLeu 220
Db |||||
QY 585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCCIG 644
Db |||||
QY 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db |||||
QY 645 GTCATGCTGGAGATGACCCCTCATCAGGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Db |||||
QY 241 SerLeuLysSerProIleThrValGluTyrArgAlaGlnSerGluSerAlaArgSerLys 260
Db |||||
QY 705 AGCCTGAAGAGCCCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Db |||||
QY 261 GlyGlyGlyGlySer 265
Db |||||
QY 756 GGCGGTGGTGGTTCC 770
Db |||||
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Search completed: June 30, 2006, 01:48:08
Job time : 539.723 secs

GenCore version 5.1.9
Copyright (c) 1993 - 2006 Bioceleration Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: June 30, 2006, 01:28:47 ; Search time 6366.34 Seconds
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4520.078 Million cell updates/sec

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Perfect score: 1572
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Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 6366136 seqs, 31973710525 residues
Total number of hits satisfying chosen parameters: 12732272

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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-USER=US10048116 @CGN 1.1 7274 @runat.29062006.093311.10139 -NCPU=6 -ICPU=3
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-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

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15: gb_ba:*

ALIGNMENTS

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DEFINITION Sequence 2 from Patent WO0109194.
ACCESSION AX081281
VERSION AX081281.1 GI:13170131
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source

synthetic construct
other sequences; artificial sequences.

1
Glaichenhaus, N. and Malherbe, L.
Recombinant proteins and molecular complexes derived therefrom,
analogous to molecules involved in immune responses
Patent: WO 0109194-A 2 08-FEB-2001;
CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)

Location/Qualifiers
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SUMMARIES

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2	1255.5	79.9	893	2 AR047947	Sequence
3	1161.5	73.9	4724	2 AR199666	Sequence

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.


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Qy      201 ThrValGlyValSerSerThrGlnLeuLeuIleArgAsnGlyAspTrrPheGlnValIleu 220
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Qy      241 SerLeuLysSerProIleThrValGluTrrPArgAlaGlnSerGluSerAlaArgSerLys 260
Db      766 AGCCTGAAGAGCCCATCATCTGTGGAGTGGAGGCGACAGTCCGAGTCTGCCCGGAGCAAG 825

RESULT 3
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LOCUS      AR199666              4724 bp      DNA      linear      PAT 20-APR-2002
DEFINITION Sequence 8 from patent US 6355479.
ACCESSION AR199666
VERSION    AR199666.1 GI:20249740
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 4724)
AUTHORS   Webb,S.R., Winqvist,O., Karlsson,L., Jackson,M.R. and Peterson,P.A.
TITLE     MHC class II antigen-presenting systems and methods for activating
          CD4+ T cells
JOURNAL    Patent: US 6355479-A 8 12-MAR-2002;
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Best Local Similarity: 85.4%      Mismatches:  1
Query Match:    73.9%      Indels:      37
DB:             2           Gaps:          2

US-10-048-116B-6_COPY_1_300 (1-300) x AR199666 (1-4724)
Qy      1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValIleu 20
Db      451 ATGGCTCTGAGATCCCGACGCTCTCTCTTCAGTCTGTGTGGTGGTGTGTGTGTGTGTGTG 510
Qy      21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      511 AGCAGCCAGGAGCTAGAGGCGGAAAC----- 537
Qy      41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db      537 ----- 537

Qy      61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrTyrThrAsn 80
Db      538 -----TCCGAAAGGCATTTCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 588
Qy      81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db      589 GGCAGCGCAGCATACGGCTCGTGACCATATACATCTACACCCGGGAGGAGTACGTGCGC 648
Qy      101 TyrAspSerAspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db      649 TACGACAGCGACGTGGCGGAGTACCGCGGTGTACCGAGCTGGGGCGCGCAGACGCGCAG 708
Qy      121 TyrTrpAsnSerGlnProGluIleLeuGluArgThrArgAlaGluValValAspThrAlaCys 140
Db      709 TACTGGNACAGCCCGCGAGATCTCTGGAGCGGAACCGGGGCGGAGGTGGACACGGCGGTGC 768
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Qy      141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgLeuGluGlnProAsn 160
Db      769 AGACACAACATACGAGGGGCGGACAGCACCTCCCTGCGGCGCTTTGAACAGCCCAAT 828
Qy      161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180
Db      829 ATCCCATCTCCCTGTCCAGACAGAGGCCCTCAACCAACCACTCTGGTCTGTGTCTG 888
Qy      181 ValThrAspPheTyrProAlaLysIleLysValArgTrrPheArgAsnGlyGlnGluGlu 200
Db      889 GTGACAGATTCTACCCAGCCCAAGATCAAAGTGGCTGGTTCAGGAATGGCCAGGAGGAG 948
Qy      201 ThrValGlyValSerSerThrGlnLeuLeuIleArgAsnGlyAspTrrPheGlnValIleu 220
Db      949 ACAGTGGGGTCTCATCCACACAGCTATTATTAGGAATGGGACTGGACCTTCAGGTCCTG 1008
Qy      221 ValMetLeuGluMetThrProHisGlnGlyValTyrThrCysHisValGluHisPro 240
Db      1009 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 1068
Qy      241 SerLeuLysSerProIleThrValGluTrrPArgAlaGlnSerGluSerAlaArgSerLys 260
Db      1069 AGCCTGAAGAGCCCATCATCTGTGGAGTGGAGGCGACAGTCCGAGTCTGCCCGGAGCAAG 1128
Qy      261 -----GlyGlyGlyGly 264
Db      1129 ATGTTGAGCGCATCGGGGCG 1149

RESULT 4
AR047957
LOCUS      AR047957              1013 bp      DNA      linear      PAT 29-SEP-1999
DEFINITION Sequence 38 from patent US 5820866.
ACCESSION AR047957
VERSION    AR047957.1 GI:5970300
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 1013)
AUTHORS   Kappler,J.W. and Marrack,P.
TITLE     Product and process for T cell regulation
JOURNAL    Patent: US 5820866-A 38 13-OCT-1998;
FEATURES   Location/Qualifiers
            source          1..1013
                        /organism="unknown"
                        /mol_type="unassigned DNA"

ORIGIN
Alignment Scores:
Pred. No.:      8.51e-112      Length:      1013
Score:          1154.00      Matches:      231
Percent Similarity: 86.8%      Conservative: 5
Best Local Similarity: 84.9%      Mismatches:  22
Query Match:    73.4%      Indels:      14
DB:             2           Gaps:          4

US-10-048-116B-6_COPY_1_300 (1-300) x AR047957 (1-1013)
Qy      1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValIleu 20
Db      63 ATGGCTCTGAGATCCCGACGCTCTCTCTCTCGCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 122
Qy      21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db      123 AGCAGCCAGGAGCTAGAGGCGGAGACTCC-----GAAGCTACTTTGAGGCTCAG 173
Qy      41 -----IleValValSerGlySerTrpAspGlyGlyGlySerLeuVal 55
Db      174 GGTGCAGCTGCCCAACATTTGCTGTGCAAGGCTGGAGGTGGTGGATCCGGTGA----- 227
Qy      56 ProArgGlySerGlyGlyGlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlu 75
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ORIGIN

Alignment Scores:
 Pred. No.: 1.13e-110 Length: 1382
 Score: 1145.00 Matches: 227
 Percent Similarity: 87.2% Conservatives: 4
 Best Local Similarity: 85.7% Mismatches: 24
 Query Match: 72.8% Indels: 10
 DB: 2 Gaps: 3

US-10-048-116B-6_COPY_1_300 (1-300) x AR175097 (1-1382)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20
 Db 6 ATGGCTCTGCAGATCCCGAGCCTCTCTCTCAGCTGCTGGTGGTCTCATGGTGTG 65

Qy 21 SerSerProGlyThrGluGlyGlyAAsnSerIleCysPheSerProSerLeuGluHisPro 40
 Db 66 AGCAGCCCAAGACCTTAAGTATCTCTCAGGCTGTTCACGCTGCTCAGCTGAA----- 119

Qy 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
 Db 120 ATCAACGAAGCTGCTGTAGCGAGGGGGCGGAAGC-----GGCGGA 164

Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrThrAsn 80
 Db 165 GGGGGAACCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTGTACTACCAAC 224

Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGlyTyrValArg 100
 Db 225 GGGACGCGAGCATACGGCTCGTACCCAGATACATCTACACCGGGAGGAGTAGTGGCG 284

Qy 101 TyrAspSerAspValGlyGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
 Db 285 GTGCCATCTCCTGTCTCAGACAGAGGCGCTCAACACCAACACACTCTGCTGTTCG 524

Qy 181 ValThrAspPheTyrProAlaLysIleIleValArgTrpPheArgAsnGlyGlnGluGlu 200
 Db 525 GTGACAGATTTCCTACCCAGCCAGATCAAGTGGCTGGTTTCAGGAATGGCCAGGGAG 584

Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220
 Db 585 ACAGTGGGGTCTCATCCACAGCTTATTAGGAATGGCTGGTTTCAGGAATGGCCAGGGAG 644

Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
 Db 705 AGCCTGAAGAGCCCCATCTCTGTGGAGTGG-----ACTAGTGTGGCGGTGGCAGC 755

Qy 261 GlyGlyGlyGlySer 265
 Db 756 GCGGGTGGTGGTTCC 770

RESULT 7

CS079301 1382 bp DNA linear PAT 06-MAY-2005
 LOCUS CS079301
 DEFINITION Sequence 123 from Patent EP1526141.
 ACCESSION CS079301

VERSION CS079301.1 GI:63093743

KEYWORDS .
 SOURCE unidentified
 ORGANISM unidentified
 unclassified sequences.

REFERENCE 1

AUTHORS Rhode,P.R., Jiao,J.A., Burkhardt,M. and Wong,H.C.

TITLE MHC complexes and uses thereof

JOURNAL Patent; EP 1526141-A 123 27-APR-2005;

Altor BioScience Corporation (US)

FEATURES

Location/Qualifiers

1..1382

source

/organism="unidentified"

/mol_type="unassigned DNA"

/db_xref="taxon:32644"

ORIGIN

Alignment Scores:

Pred. No.: 1.13e-110 Length: 1382

Score: 1145.00 Matches: 227

Percent Similarity: 87.2% Conservatives: 4

Best Local Similarity: 85.7% Mismatches: 24

Query Match: 72.8% Indels: 10

DB: 2 Gaps: 3

US-10-048-116B-6_COPY_1_300 (1-300) x CS079301 (1-1382)

Qy 1 MetAlaLeuGlnIleProSerLeuLeuSerAlaAlaValValValLeuMetValLeu 20

Db 6 ATGGCTCTGCAGATCCCGAGCCTCTCTCTCAGCTGCTGGTGGTCTCATGGTGTG 65

Qy 21 SerSerProGlyThrGluGlyGlyAAsnSerIleCysPheSerProSerLeuGluHisPro 40

Db 66 AGCAGCCCAAGACCTTAAGTATCTCTCAGGCTGTTCACGCTGCTCAGCTGAA----- 119

Qy 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60

Db 120 ATCAACGAAGCTGCTGTAGCGAGGGGGCGGAAGC-----GGCGGA 164

Qy 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGlyCysTyrThrAsn 80

Db 165 GGGGGAACCTCCGAAAGGCATTTCTGTGTCAGTTCAAGGGCGAGTGTACTACCAAC 224

Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGlyTyrValArg 100

Db 225 GGGACGCGAGCATACGGCTCGTACCCAGATACATCTACACCGGGAGGAGTAGTGGCG 284

Qy 101 TyrAspSerAspValGlyGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120

Db 285 GTGCCATCTCCTGTCTCAGACAGAGGCGCTCAACACCAACACACTCTGCTGTTCG 524

Qy 181 ValThrAspPheTyrProAlaLysIleIleValArgTrpPheArgAsnGlyGlnGluGlu 200

Db 525 GTGACAGATTTCCTACCCAGCCAGATCAAGTGGCTGGTTTCAGGAATGGCCAGGGAG 584

Qy 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTrpPheGlnValLeu 220

Db 585 ACAGTGGGGTCTCATCCACAGCTTATTAGGAATGGCTGGTTTCAGGAATGGCCAGGGAG 644

Qy 221 ValMetLeuGluMetThrProHisGlnGlyGlyValTyrThrCysHisValGluHisPro 240

Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGCTTACACCTGCCATGTGGAGCATCCC 704

Qy 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260

Db 705 AGCCTGAAGAGCCCCATCTCTGTGGAGTGG-----ACTAGTGTGGCGGTGGCAGC 755

Qy 261 GlyGlyGlyGlySer 265

Db 756 GCGGGTGGTGGTTCC 770

Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
QY 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
QY 261 GlyGlyGlyGlySer 265
Db 756 GCGGTGGTGGTTCC 770
RESULT 13
AR033963
LOCUS AR033963 1508 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 122 from patent US 5869270.
ACCESSION AR033963
VERSION AR033963.1 GI:5949568
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1508)
AUTHORS Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
TITLE Single chain MHC complexes and uses thereof
JOURNAL Patent: US 5869270-A 122 09-FEB-1999;
FEATURES Location/Qualifiers
source
1..1508
/organism="unknown"
/mol_type="unassigned DNA"
ORIGIN
Alignment Scores:
Pred. No.: 1.26e-110 Length: 1508
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 72.8% Indels: 10
DB: 2 Gaps: 3
US-10-048-116b-6_COPY_1_300 (1-300) x AR033963 (1-1508)
QY 1 MetalaleuGlnIleProSerLeuLeuLeuSerAlaAlaValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGTTCAGCTGTTCAGCTGAA-----119
QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGACCTTAAGTATCTCTCAGCTGTTCAGCTGTTCAGCTGAA-----119
QY 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAAGCTGTGCTGCTAGCGAGGGGGCGGAAGC-----GGCGGA 164
QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrThrAsn 80
Db 165 GGGGGAACCTCCGAAGCATTTCTGTCAGTTCAGGCGGAGGTGACTACACCAAC 224
QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGACGCAGCGCATACGCTGCTGCTAGCGAGGGGGCGGAAGC-----GGCGGA 164
QY 101 TyrAspSerAspValGlyTyrArgAlaValThrGluLeuGlyArgProAspAlaGlu 120
Db 285 TACGACAGCGAGCTGGGCGAGTACCGCGGTGACCGAGCTGGGGCGGCCAGCGCGAG 344
QY 121 TyrTrpAsnSerGlnProGluIleGluArgThrArgAlaGluValAspThrAlaCys 140
Db 345 TACTGGAAACACCGACCGAGATCTTGAGGGAACCGCGCGGAGGTGGACCGCGGTGC 404
QY 141 ArgHisAsnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAsn 160
Db 405 AGACACAACATACGAGGGCGGAGACCGACACTCTCTCGCGCGCTTGAACGCCCAAT 464
QY 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAsnHisHisAsnThrLeuValCysSer 180

Db 465 GTCGCATCTCCCTGTCCAGGACAGAGCCCTCAACACCACCAACACTCTGGTCTGTTCG 524
QY 181 ValThrAspPheTyrProAlaLysIleLysValArgTTPheArgAsnGlyGlnGlu 200
Db 525 GTGACAGATTTCTACCCAGCCAAGATCAAGTGCCTGTTCCAGGAATGCCAGGAGGAG 584
QY 201 ThrValGlyValSerSerThrGlnLeuIleArgAsnGlyAspTyrThrPheGlnValLeu 220
Db 585 ACAGTGGGGTCTCATCCACACAGCTTATTAGGAATGGGACTGGACCTTCAGGTCTCTG 644
QY 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCATGTGGAGCATCCC 704
QY 241 SerLeuLysSerProIleThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
QY 261 GlyGlyGlyGlySer 265
Db 756 GCGGTGGTGGTTCC 770
RESULT 14
AR152030
LOCUS AR152030 1508 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 24 from patent US 6232445.
ACCESSION AR152030
VERSION AR152030.1 GI:15118080
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1508)
AUTHORS Rhode,P.R., Acevedo,J., Burkhardt,M., Jiao,J.-a. and Wong,H.C.
TITLE Soluble MHC complexes and methods of use thereof
JOURNAL Patent: US 6232445-A 24 15-MAY-2001;
FEATURES Location/Qualifiers
source
1..1508
/organism="unknown"
/mol_type="unassigned DNA"
ORIGIN
Alignment Scores:
Pred. No.: 1.26e-110 Length: 1508
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 72.8% Indels: 10
DB: 2 Gaps: 3
US-10-048-116b-6_COPY_1_300 (1-300) x AR152030 (1-1508)
QY 1 MetalaleuGlnIleProSerLeuLeuLeuSerAlaAlaValValLeuMetValLeu 20
Db 6 ATGGCTCTGCAGATCCCGAGCTCTCTCTCAGCTGTTCAGCTGTTCAGCTGAA-----65
QY 21 SerSerProGlyThrGluGlyGlyAsnSerIleCysPheSerProSerLeuGluHisPro 40
Db 66 AGCAGCCCAAGACCTTAAGTATCTCTCAGCTGTTCAGCTGTTCAGCTGAA-----119
QY 41 IleValValSerGlySerTrpAspGlyGlyGlySerLeuValProArgGlySerGly 60
Db 120 ATCAACGAAGCTGTGCTGCTAGCGAGGGGGCGGAAGC-----GGCGGA 164
QY 61 GlyGlyGlySerGluArgHisPheValValGlnPheLysGlyGluCysTyrThrAsn 80
Db 165 GGGGGAACCTCCGAAGCATTTCTGTCAGTTCAGGCGGAGGTGACTACACCAAC 224
QY 81 GlyThrGlnArgIleArgLeuValThrArgTyrIleTyrAsnArgGluGluTyrValArg 100
Db 225 GGGACGCAGCGCATACGCTGCTGCTAGCGAGGGGGCGGAAGC-----GGCGGA 164

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Qy 101 TyrAaspSerAapValGlyGluTyrArgAlaValThrGluLeuGlyArgProAaspAlaGlu 120
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Db 285 TAGACAGCAGCGTGGCGGAGTACCGCGCGGTGACCGAGCTGGGGCGCCAGACGCCGAG 344
Qy 121 TyrTrpAasnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAaspThrAlaCys 140
    |||
Db 345 TACTGGAACAGCCAGCCGGAGATCCTGGAGCGAACCGCGCGCGAGGTGGACACGGCGTGC 404
Qy 141 ArgHisAenTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAasn 160
    |||
Db 405 AGACACAACTACGAGGGGCGGAGACAGCACCTCCCTGCGGGCGCTTGAACAGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAasnHisAenThrLeuValCysSer 180
    |||
Db 465 GTGCCATCTCCCTGTCAGACAGAGCGCCCTCAACCAACACACACTCTGGTCTGTTCG 524
Qy 181 ValThrAaspPheTyrProAlaLysIleLysValArgTrpPheArgAasnGlyGlnGluGlu 200
    |||
Db 525 GTGACAGATTCTACCCAGCCAGATCAAAGTGGCTGGTTTCAGGAATGGCCAGGAGGAG 584
Qy 201 ThrValGlyValSerSerThrGlnLeuLeuArgAasnGlyAaspThrPheGlnValLeu 220
    |||
Db 585 ACAGTGGGGTCTCATCACACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTCG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
    |||
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy 241 SerLeuLysSerProfileThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
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Db 756 GCGGTGGTGGTTCC 770
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RESULT 15

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DEFINITION Sequence 122 from patent US 6309645.
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ACCESSION ARI175096
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VERSION ARI175096.1 GI:17916395
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KEYWORDS
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SOURCE Unknown.
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ORGANISM Unknown.
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REFERENCE 1. (bases 1 to 1508)
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AUTHORS Rhode,P.R., Jiao,J.-A., Burkhardt,M. and Wong,H.C.
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TITLE MHC molecules and uses thereof
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JOURNAL Patent: US 6309645-A 122 30-OCT-2001;
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ORIGIN

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Alignment Scores:
Pred. No.: 1.26e-110 Length: 1508
Score: 1145.00 Matches: 227
Percent Similarity: 87.2% Conservative: 4
Best Local Similarity: 85.7% Mismatches: 24
Query Match: 72.8% Indels: 10
DB: 2 Gaps: 3
```

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US-10-048-116B-6_COPY_1_300 (1-300) x ARI175096 (1-1508)
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Qy 21 SerSerProGlyThrGluGlyGlyAasnIleCysPheSerProSerLeuGluHisPro 40
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Db 66 AGCAGCCCAAGGACCTTAAAGTATCTCTCAGGCTGTTCCAGCTGCTCAGCTGAA----- 119
```

Search completed: June 30, 2006, 05:17:57

Job time : 6369.34 secs

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    |||
Db 165 GGGGGAAACTCCGAAAGGCATTTTCGTGGTCCAGTTCAAGGGCGAGTGCTACTACACCAAC 224
Qy 81 GlyThrGlnArgIleArgLeuValThrArgTyrTyrIleTyrAasnArgGluGluTyrValArg 100
    |||
Db 225 GGGACACAGCGCATACGGCTCGTGACCAATCATCTACAACCGGGAGGAGTAGTGTCGC 284
Qy 101 TyrAaspSerAaspValGlyGluTyrArgAlaValThrGluLeuGlyArgProAaspAlaGlu 120
    |||
Db 285 TAGCACAGGACGAGTGGCGGAGTACCGCGCGGTGACCGAGTGGGGCGCCAGCGCGAG 344
Qy 121 TyrTrpAasnSerGlnProGluLeuLeuGluArgThrArgAlaGluValAaspThrAlaCys 140
    |||
Db 345 TACTGGAACAGCCAGCCGAGATCCTTGGAGCAACCGCGCGCGAGGTGGACACCGCGTGC 404
Qy 141 ArgHisAasnTyrGluGlyProGluThrSerThrSerLeuArgArgLeuGluGlnProAasn 160
    |||
Db 405 AGACACAACTACGAGGGGCGGAGACAGCACCTCCCTGCGGGCGCTTGAACAGCCCAAT 464
Qy 161 ValAlaIleSerLeuSerArgThrGluAlaLeuAasnHisAenThrLeuValCysSer 180
    |||
Db 465 GTGCCATCTCCCTGTCAGGACAGAGCGCCCTCAACCAACCAACACTCTGGTCTGTTCG 524
Qy 181 ValThrAaspPheTyrProAlaLysIleLysValArgTrpPheArgAasnGlyGlnGluGlu 200
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Qy 201 ThrValGlyValSerSerThrGlnLeuLeuArgAasnGlyAaspThrPheGlnValLeu 220
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Db 585 ACAGTGGGGTCTCATCCACAGCTTATTAGGAATGGGACTGGACCTTCCAGGTCTCG 644
Qy 221 ValMetLeuGluMetThrProHisGlnGlyGluValTyrThrCysHisValGluHisPro 240
    |||
Db 645 GTCATGCTGGAGATGACCCCTCATCAGGAGAGGTCTACACCTGCCATGTGGAGCATCCC 704
Qy 241 SerLeuLysSerProfileThrValGluTrpArgAlaGlnSerGluSerAlaArgSerLys 260
    |||
Db 705 AGCCTGAAGAGCCCATCACTGTGGAGTGG-----ACTAGTGGTGGCGGTGGCAGC 755
Qy 261 GlyGlyGlyGlySer 265
Db 756 GCGGTGGTGGTTCC 770
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